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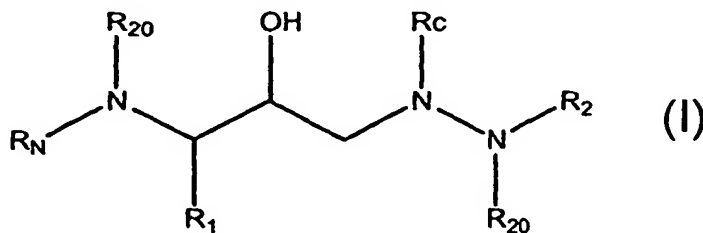
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(54) Title: AZA HYDROXYLATED ETHYL AMINE COMPOUNDS

(57) Abstract: Disclosed are
compounds of formula (I) and
pharmaceutically acceptable salts and
esters thereof, useful in treating and/or
preventing Alzheimer's disease and
other similar diseases, wherein R_N,
R_C, R₁, R₂ and R₂₀ are defined herein.
These compounds include inhibitors
of the beta-secretase enzyme that are
useful in the treatment of Alzheimer's
disease and other diseases characterized
by deposition of A beta peptide in
a mammal. The compounds of the
invention are useful in pharmaceutical

compositions and methods of treatment to reduce A beta peptide formation.

WO 02/094768 A2

AZA HYDROXYLATED ETHYL AMINE COMPOUNDS

(Case No. 02-064-C)

5

Background of the Invention**Field of the Invention**

This invention relates to aza hydroxylated ethyl amine derivatives and more specifically to such compounds that are useful in the treatment and/or prevention of Alzheimer's disease and similar diseases. More specifically the invention relates to substituted aza hydroxy ethyl amines that are capable of inhibiting beta-secretase, an enzyme that cleaves amyloid precursor protein to produce amyloid beta peptide (A beta), a major component of the amyloid plaques found in the brains of Alzheimer's sufferers.

Description of Related Art

Alzheimer's disease (AD) is a progressive degenerative disease of the brain primarily associated with aging. Clinical presentation of AD is characterized by loss of memory, cognition, reasoning, judgment, and orientation. As the disease progresses, motor, sensory, and linguistic abilities are also affected until there is global impairment of multiple cognitive functions. These cognitive losses occur gradually, but typically lead to severe impairment and eventual death in the range of four to twelve years.

Alzheimer's disease is characterized by two major pathologic observations in the brain: neurofibrillary tangles and beta amyloid (or neuritic) plaques, comprised predominantly of an aggregate of a peptide fragment known as A beta. Individuals with AD exhibit characteristic beta-amyloid deposits in the brain (beta amyloid plaques) and in cerebral blood vessels (beta amyloid angiopathy) as well as neurofibrillary tangles. Neurofibrillary tangles occur not only in Alzheimer's disease but also in other dementia-inducing disorders. On

autopsy, large numbers of these lesions are generally found in areas of the human brain important for memory and cognition.

Smaller numbers of these lesions in a more restricted anatomical distribution are found in the brains of most aged humans who do not have clinical AD. Amyloidogenic plaques and vascular amyloid angiopathy also characterize the brains of individuals with Trisomy 21 (Down's Syndrome), Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type (HCHWA-D), and other neurodegenerative disorders. Beta-amyloid is a defining feature of AD, now believed to be a causative precursor or factor in the development of disease. Deposition of A beta in areas of the brain responsible for cognitive activities is a major factor in the development of AD. Beta-amyloid plaques are predominantly composed of amyloid beta peptide (A beta, also sometimes designated betaA4). A beta peptide is derived by proteolysis of the amyloid precursor protein (APP) and is comprised of 39-42 amino acids. Several proteases called secretases are involved in the processing of APP.

Cleavage of APP at the N-terminus of the A beta peptide by beta-secretase and at the C-terminus by one or more gamma-secretases constitutes the beta-amyloidogenic pathway, i.e. the pathway by which A beta is formed. Cleavage of APP by alpha-secretase produces alpha-sAPP, a secreted form of APP that does not result in beta-amyloid plaque formation. This alternate pathway precludes the formation of A beta peptide. A description of the proteolytic processing fragments of APP is found, for example, in U.S. Patent Nos. 5,441,870; 5,721,130; and 5,942,400.

An aspartyl protease has been identified as the enzyme responsible for processing of APP at the beta-secretase cleavage site. The beta-secretase enzyme has been disclosed using varied nomenclature, including BACE, Asp, and Memapsin. See, for example, Sindha et al., 1999, *Nature* 402:537-554 (p501) and published PCT application WO00/17369.

Several lines of evidence indicate that progressive cerebral deposition of beta-amyloid peptide (A beta) plays a seminal role in the pathogenesis of AD and can precede cognitive symptoms by years or decades. See, for example, Selkoe, 1991, *Neuron* 6:487. Release of A beta from neuronal cells grown in culture and the presence of A beta in cerebrospinal fluid (CSF) of both normal individuals and AD patients has been demonstrated. See, for example, Seubert et al., 1992, *Nature* 359:325-327.

It has been proposed that A beta peptide accumulates as a result of APP processing by beta-secretase, thus inhibition of this enzyme's activity is desirable for the treatment of AD. *In vivo* processing of APP at the beta-secretase cleavage site is thought to be a rate-limiting step in A beta production, and is thus a therapeutic target for the treatment of AD. See for example, Sabbagh, M., et al., 1997, *Alz. Dis. Rev.* 3, 1-19.

BACE1 knockout mice fail to produce A beta, and present a normal phenotype. When crossed with transgenic mice that over express APP, the progeny show reduced amounts of A beta in brain extracts as compared with control animals (Luo et al., 2001 *Nature Neuroscience* 4:231-232). This evidence further supports the proposal that inhibition of beta-secretase activity and reduction of A beta in the brain provides a therapeutic method for the treatment of AD and other beta amyloid disorders.

At present there are no effective treatments for halting, preventing, or reversing the progression of Alzheimer's disease. Therefore, there is an urgent need for pharmaceutical agents capable of slowing the progression of Alzheimer's disease and/or preventing it in the first place.

Compounds that are effective inhibitors of beta-secretase, that inhibit beta-secretase-mediated cleavage of APP, that are effective inhibitors of A beta production, and/or are effective to reduce amyloid beta deposits or plaques, are needed for the treatment and prevention of disease characterized by amyloid beta deposits or plaques, such as AD.

Various pharmaceutical agents have been proposed for the treatment of Alzheimer's disease but without any real success.

At present there are no effective treatments for halting, preventing, or reversing the progression of Alzheimer's disease.

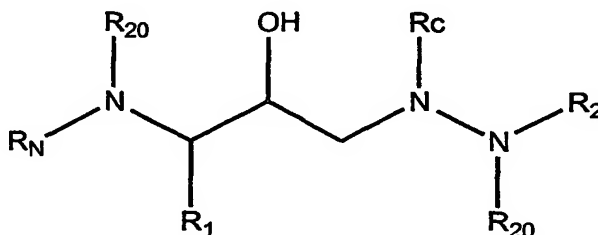
5 There is an urgent need for compounds capable of slowing A-beta peptide production and/or deposition in the brain, which presents a therapeutic approach to treatment of Alzheimer's disease.

Summary of the Invention

10 The invention provides compounds of the formula below, pharmaceutical compositions containing the compounds, and methods useful in the treatment of Alzheimer's disease. More specifically, it provides compounds that are capable of inhibiting beta-secretase, an enzyme that cleaves amyloid precursor protein to produce A beta peptide, a major component of the amyloid plaques found in the brains of Alzheimer's sufferers.

Accordingly, in a broad aspect, the invention is provides towards compounds of formula I:

20



(I)

25 and pharmaceutically acceptable salts or esters thereof, where R_c is

(I) -C₁-C₁₀ alkyl optionally substituted with one, two or three groups independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, -NR_{1-a}R_{1-b}, -OC(=O)NR_{1-a}R_{1-b}, -S(=O)₀₋₂R_{1-a}, -NR_{1-a}C(=O)NR_{1-a}R_{1-b}, -C(=O)NR_{1-a}R_{1-b}, and -S(=O)₂NR_{1-a}R_{1-b} wherein

R_{1-a} and R_{1-b} at each occurrence are independently H or C_1-C_6 alkyl,

(II) $-(CH_2)_{0-3}-(C_3-C_8)$ cycloalkyl where cycloalkyl can be optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, halogen, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O$ -phenyl, $-CO_2H$, $-CO_2-(C_1-C_4 \text{ alkyl})$, and $-NR_{1-a}R_{1-b}$

(III) $-(CR_{C-x}R_{C-y})_{0-4}-R_{C-aryl}$ where R_{C-x} and R_{C-y} are independently selected from the group consisting of

10 $-H$,
 C_1-C_4 alkyl optionally substituted with 1 or 2 $-OH$,
 C_1-C_4 alkoxy optionally substituted with 1, 2, or 3
halogen,
 $-(CH_2)_{0-4}-C_3-C_8$ cycloalkyl,
15 C_2-C_6 alkenyl containing one or two double bonds,
 C_2-C_6 alkynyl containing one or two triple bonds, and
phenyl,

or

R_{C-x} and R_{C-y} are taken together with the carbon to
20 which they are attached to form a carbocycle of three, four, five, six or seven carbon atoms, where one carbon atom is optionally replaced by a group selected from $-O-$, $-S-$, $-SO_2-$, $-NR_{N-2}-$ and R_{C-aryl} , wherein

R_{C-aryl} at each occurrence is independently phenyl;
25 naphthyl; tetralinyl; indanyl; dihydronaphthyl; or 6,7,8,9-tetrahydro-5H-benzo[a]cycloheptenyl, each of which is optionally substituted with 1, 2, or 3 groups that are independently:

(1) C_1-C_6 alkyl, optionally substituted with one, two or three substituents selected from the group
30 consisting of C_1-C_3 alkyl, halogen, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(2) $-OH$,

(3) $-NO_2$,

(4) halogen,

35 (5) $-CO_2H$,

(6) $-\text{C}\equiv\text{N}$,

(7) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$ where

$\text{R}_{\text{N}-2}$ and $\text{R}_{\text{N}-3}$ are independently selected from the group consisting of:

5

(a) $-\text{H}$,

(b) $-\text{C}_1-\text{C}_6$ alkyl optionally substituted with one substituent selected from the group consisting of:

(i) $-\text{OH}$, and

(ii) $-\text{NH}_2$,

10

(c) $-\text{C}_1-\text{C}_6$ alkyl optionally substituted with 1, 2, or 3 groups that are independently $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, or OH ,

(d) $-\text{C}_3-\text{C}_7$ cycloalkyl,

(e) $-(\text{C}_1-\text{C}_2 \text{ alkyl})-(\text{C}_3-\text{C}_7 \text{ cycloalkyl})$,

15

(f) $-(\text{C}_1-\text{C}_6 \text{ alkyl})-\text{O}-(\text{C}_1-\text{C}_3 \text{ alkyl})$,

(g) $-\text{C}_2-\text{C}_6$ alkenyl

(h) $-\text{C}_2-\text{C}_6$ alkynyl

(i) $-\text{C}_1-\text{C}_6$ alkyl chain with one double bond and one triple bond,

20

(j) $-\text{R}_{1-\text{aryl}}$ wherein $\text{R}_{1-\text{aryl}}$ at each occurrence is independently phenyl, naphthyl, indanyl, indenyl, dihydronaphthyl, or tetralinyl each of which is optionally substituted with 1, 2, 3, or 4 groups that are independently:

25

(i) C_1-C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{NR}_{1-\text{a}}\text{R}_{1-\text{b}}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, and C_1-C_3 alkoxy,

30

(ii) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, and $-\text{NR}_{1-\text{a}}\text{R}_{1-\text{b}}$,

(iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently

selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(iv) -F, Cl, -Br and -I,

(v) -C₁-C₆ alkoxy optionally

5 substituted with 1, 2, or 3 -F,

(vi) -NR_{N-2}R_{N-3},

(vii) -OH,

(viii) -C≡N,

(ix) C₃-C₇ cycloalkyl, optionally

10 substituted with 1, 2, or 3 groups that are selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(x) -CO-(C₁-C₄ alkyl),

(xi) -SO₂-NR_{1-a}R_{1-b},

15

(xii) -CO-NR_{1-a}R_{1-b}, or

(xiii) -SO₂-(C₁-C₄ alkyl),

(k) -R_{1-heteroaryl} wherein R_{1-heteroaryl} at

each occurrence is independently selected from the group consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl,

20 indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl,

isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl,

imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl,

indolizinyl, indazolyl, benzothiazolyl, benzimidazolyl,

benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl,

25 thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl,

imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl,

carbazolyl, beta-carbolinyl, isochromanyl, chromanyl,

tetrahydroisoquinolinyl, isoindolinyl,

isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl,

30 isobenzothienyl, benzoxazolyl, pyridopyridinyl,

benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl,

benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl,

pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,

dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,

dihydrobenzothiazinyl, benzopyranyl, benzothiopyranyl,
coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-
oxide, tetrahydroquinolinyl, dihydroquinolinyl,
dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
5 dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl,
benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,
indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide,
quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-
10 oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
thiazolyl N-oxide, indolizinyll N-oxide, indazolyl N-oxide,
benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and
15 benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is
optionally substituted with 1, 2, 3, or 4 groups that are
independently:

(i) C_1 - C_6 alkyl optionally
20 substituted with 1, 2, or 3 groups independently selected from
the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH,
- $NR_{1-a}R_{1-b}$, -C \equiv N, -CF $_3$, and C_1 - C_3 alkoxy,

(ii) C_2 - C_6 alkenyl optionally
substituted with 1, 2, or 3 groups that are independently -F, -
25 Cl, -OH, -SH, -C \equiv N, -CF $_3$, C_1 - C_3 alkoxy, and - $NR_{1-a}R_{1-b}$,

(iii) C_2 - C_6 alkynyl
optionally substituted with 1, 2, or 3 groups that are
independently selected from the group consisting of -F, -Cl,
-OH, -SH, -C \equiv N, -CF $_3$, C_1 - C_3 alkoxy, and - $NR_{1-a}R_{1-b}$,

30 (iv) -F, -Cl, -Br and -I,

(v) - C_1 - C_6 alkoxy optionally
substituted with one, two, or three -F,

(vi) -(CH $_2$) $_{0-4}$ - $NR_{N-2}R_{N-3}$,

(vii) -OH,

(viii) $-C\equiv N$,
 (ix) $(CH_2)_{0-4}-C_3-C_7$ cycloalkyl,
 optionally substituted with 1, 2, or 3 groups that are
 independently selected from the group consisting of $-F$, $-Cl$,
 5 $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(x) $(CH_2)_{0-4}-CO-(C_1-C_6 \text{ alkyl})$,
 (xi) $(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,
 (xii) $(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,
 (xiii) $(CH_2)_{0-4}-SO_2-(C_1-C_6$

10 $\text{alkyl})$,

(xiv) $(CH_2)_{0-4}-N(R_{N-2})-SO_2-$,

and

(xv) $(CH_2)_{0-4}-N(R_{N-2})-C(O)-$,

- 15 (8) $-(CH_2)_{0-4}-CO-(C_1-C_{12} \text{ alkyl})$,
 (9) $-(CH_2)_{0-4}-CO-(C_2-C_{12} \text{ alkenyl})$,
 (10) $-(CH_2)_{0-4}-CO-(C_2-C_{12} \text{ alkynyl})$,
 (11) $-(CH_2)_{0-4}-CO-(CH_2)_{0-4} (C_3-C_7 \text{ cycloalkyl})$,
 (12) $-(CH_2)_{0-4}-CO-R_1\text{-aryl}$,
 (13) $-(CH_2)_{0-4}-CO-R_1\text{-heteroaryl}$,
 20 (14) $-(CH_2)_{0-4}-CO-R_1\text{-heterocycle}$ wherein

$R_1\text{-heterocycle}$ at each occurrence is
 independently selected from the group consisting of morpholinyl,
 thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-
 dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl,
 25 tetrahydropyranyl, piperidinyl, tetrahydrofuranyl,
 tetrahydrothienyl, homopiperidinyl, homomorpholinyl,
 homothiomorpholinyl, homothiomorpholinyl S,S-dioxide,
 oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl,
 dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl,
 30 dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide,
 tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the $R_1\text{-heterocycle}$ group is bonded by
 any atom of the parent $R_1\text{-heterocycle}$ group substituted by hydrogen
 such that the new bond to the $R_1\text{-heterocycle}$ group replaces the

hydrogen atom and its bond, where heterocycle is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(a) C_1-C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, halogen, -OH, -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

(b) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(c) C_2-C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(d) halogen,

(e) C_1-C_6 alkoxy,

(f) $-C_1-C_6$ alkoxy optionally substituted with one, two, or three -F,

(g) $-NR_{N-2}R_{N-3}$,

(h) -OH,

(i) $-C\equiv N$,

(j) $(CH_2)_{0-4}-(C_3-C_7$ cycloalkyl), optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(k) $-(CH_2)_{0-4}-CO-(C_1-C_4$ alkyl),

(l) $-(CH_2)_{0-4}-SO_2-NR_{1-a}R_{1-b}$,

(m) $-(CH_2)_{0-4}-CO-NR_{1-a}R_{1-b}$,

(n) $-(CH_2)_{0-4}-SO_2-(C_1-C_6$ alkyl), and

(o) =O,

(p) $-(CH_2)_{0-4}-N(R_{N-2})-SO_2-$

(q) $-(CH_2)_{0-4}-N(R_{N-2})-C(O)-$

(15) $-(CH_2)_{0-4}-CO-R_{N-4}$ wherein

R_{N-4} at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, pyrrolidinonyl, pyrrolyl, pyrazolyl, thienyl, pyridyl N-oxide, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl and pyrrolidinyl where each group is optionally substituted with 1, 2, 3, or 4 groups that are independently C_1-C_6 alkyl,

(16) $-(CH_2)_{0-4}-CO_2-R_{N-5}$ where

R_{N-5} at each occurrence is independently selected from the group consisting of:

- (a) C_1-C_6 alkyl,
- (b) $-(CH_2)_{0-2}-(R_{1-aryl})$,
- (c) C_2-C_6 alkenyl,
- (d) C_2-C_6 alkynyl,
- (e) C_3-C_7 cycloalkyl, and
- (f) $-(CH_2)_{0-4}-(R_{1-heteroaryl})$,

(17) $-(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$

(18) $-(CH_2)_{0-4}-SO-(C_1-C_8 \text{ alkyl})$,

(19) $-(CH_2)_{0-4}-SO_2-(C_1-C_{12} \text{ alkyl})$,

(20) $-(CH_2)_{0-4}-SO_2-(C_3-C_7 \text{ cycloalkyl})$,

(21) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO_2-R_{N-5}$,

(22) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO-N(R_{N-5})_2$,

(23) $-(CH_2)_{0-4}-N-CS-N(R_{N-5})_2$,

(24) $-(CH_2)_{0-4}-N(-H \text{ or } R_{N-5})-CO-R_{N-2}$,

(25) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,

(26) $-(CH_2)_{0-4}-R_{N-4}$,

(27) $-(CH_2)_{0-4}-O-CO-(C_1-C_6 \text{ alkyl})$,

(28) $-(CH_2)_{0-4}-O-P(O)-(OR_{100})_2$ where R_{100} is

independently H or C_1-C_4 alkyl,

(29) $-(CH_2)_{0-4}-O-CO-N(R_{N-5})_2$,

(30) $-(CH_2)_{0-4}-O-CS-N(R_{N-5})_2$,

(31) $-(CH_2)_{0-4}-O-(R_{N-5})$,

(32) $-(CH_2)_{0-4}-O-(R_{N-5})-COOH$,

(33) $-(CH_2)_{0-4}-S-(R_{N-5})$,

(34) $-(CH_2)_{0-4}-O-(C_1-C_6 \text{ alkyl})$ wherein the alkyl group is optionally substituted with one, two, three, four, or five substituents independently selected from the group consisting of F, Cl, Br, and I,

5 (35) $-(CH_2)_{0-4}-(C_3-C_8 \text{ cycloalkyl})$,

(36) C_2-C_6 alkenyl optionally substituted with C_1-C_3 alkyl, halogen, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, or $-NR_{1-a}R_{1-b}$,

10 (37) C_2-C_6 alkynyl optionally substituted with C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, or $-NR_{1-a}R_{1-b}$, and

(38) $-(CH_2)_{0-4}-N(-H \text{ or } R_{N-5})-SO_2-R_{N-2}$;

(IV) $-(CR_{C-x}R_{C-y})_{0-4}-R_{C-heteroaryl}$ wherein

$R_{C-heteroaryl}$ at each occurrence is independently selected
 15 from the group consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl, benzoisothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl,
 20 oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl,
 25 isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, hexoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl, dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
 30 dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl, coumarinyl, isocoumarinyl, chromonyl, chromanonyl, tetrahydroquinolinyl, dihydroquinolinyl, dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
 35 dihydroisocoumarinyl, isoindolinonyl,

benzodioxanyl, benzoxazolinonyl, imidazopyrazolyl,
 quinazolinonyl, pyrazopyridyl, benzooxadiazolyl,
 dihydropyrimidinonyl, dihydrobenzofuranonyl, pyridinyl-N-oxide,
 pyrrolyl N-oxide, pyrimidinyl N-oxide, pyridazinyl N-oxide,
 5 pyrazinyl N-oxide, quinolinyl N-oxide, indolyl N-oxide,
 indolinyl N-oxide, isoquinolyl N-oxide, quinazolinyl N-oxide,
 quinoxalinyl N-oxide, phthalazinyl N-oxide, imidazolyl N-oxide,
 isoxazolyl N-oxide, oxazolyl N-oxide, thiazolyl N-oxide,
 indoliziny N-oxide, indazolyl N-oxide, benzothiazolyl N-oxide,
 10 benzimidazolyl N-oxide, pyrrolyl N-oxide, oxadiazolyl N-oxide,
 thiadiazolyl N-oxide, triazolyl N-oxide, tetrazolyl N-oxide,
 benzothiopyranyl S-oxide, and benzothiopyranyl S,S-dioxide,

where the R_C-heteroaryl group is bonded by any atom of the
 parent R_C-heteroaryl group substituted by hydrogen such that the new
 15 bond to the R_C-heteroaryl group replaces the hydrogen atom and its
 bond, where heteroaryl is optionally substituted 1, 2, 3, or 4
 groups that are independently:

(1) C₁-C₆ alkyl, optionally substituted with 1, 2, or
 3 groups independently selected from the group consisting of C₁-
 20 C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy,
 and -NR_{1-a}R_{1-b},

(2) -OH,

(3) -NO₂,

(4) -F, -Cl, -Br, -I,

25 (5) -CO-OH,

(6) -C≡N,

(7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3},

(8) -(CH₂)₀₋₄-CO-(C₁-C₁₂ alkyl),

(9) -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkenyl with one, two or three
 30 double bonds),

(10) -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkynyl with one, two or three
 triple bonds),

(11) -(CH₂)₀₋₄-CO-(C₃-C₇ cycloalkyl),

(12) -(CH₂)₀₋₄-CO-R_{1-aryl} where R_{1-aryl} is as defined
 35 above,

- (13) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl}$,
 (14) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle}$,
 (15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$,
 (16) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$,
 5 (17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
 (18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1\text{-C}_8 \text{ alkyl})$,
 (19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1\text{-C}_{12} \text{ alkyl})$,
 (20) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3\text{-C}_7 \text{ cycloalkyl})$,
 (21) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or R}_{\text{N}-5})-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$,
 10 (22) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (23) $-(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (24) $-(\text{CH}_2)_{0-4}-\text{N}(-\text{H or R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2}$,
 (25) $-(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
 (26) $-(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4}$,
 15 (27) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1\text{-C}_6 \text{ alkyl})$,
 (28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$ where R_{100} is $-\text{H}$ or $\text{C}_1\text{-C}_4$
 alkyl,
 (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
 20 (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
 (32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
 (33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
 (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1\text{-C}_6 \text{ alkyl optionally substituted}$
 with one, two, three, four, or five of $-\text{F}$),
 25 (35) $\text{C}_3\text{-C}_7 \text{ cycloalkyl}$,
 (36) $\text{C}_2\text{-C}_6 \text{ alkenyl optionally substituted with C}_1\text{-C}_3$
 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1\text{-C}_3 \text{ alkoxy}$, or
 $-\text{NR}_{1-\text{a}}\text{R}_{1-\text{b}}$,
 (37) $\text{C}_2\text{-C}_6 \text{ alkynyl optionally substituted with C}_1\text{-C}_3$
 30 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1\text{-C}_3 \text{ alkoxy}$, or
 $-\text{NR}_{1-\text{a}}\text{R}_{1-\text{b}}$,
 (38) $-(\text{CH}_2)_{0-4}-\text{N}(-\text{H or R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$,
 (39) $-(\text{CH}_2)_{0-4}-(\text{C}_3\text{-C}_7 \text{ cycloalkyl})$,
 (V) $-(\text{CR}_{\text{C}-\text{x}}\text{R}_{\text{C}-\text{y}})_{0-4}-\text{R}_{\text{C}}\text{-aryl}-\text{R}_{101}-\text{R}_{\text{C}}\text{-aryl}$,
 35 (VI) $-(\text{CR}_{\text{C}-\text{x}}\text{R}_{\text{C}-\text{y}})_{0-4}-\text{R}_{\text{C}}\text{-aryl}-\text{R}_{101}-\text{R}_{\text{C}}\text{-heteroaryl}$,

- (VII) - $(CR_{C-x}R_{C-y})_{0-4}-R_C\text{-heteroaryl}-R_{101}-R_C\text{-aryl}$,
 (VIII) - $(CR_{C-x}R_{C-y})_{0-4}-R_C\text{-heteroaryl}-R_{101}-R_C\text{-heteroaryl}$,
 (IX) - $(CR_{C-x}R_{C-y})_{0-4}-R_C\text{-aryl}-R_{101}-R_1\text{-heterocycle}$,
 (X) - $(CR_{C-x}R_{C-y})_{0-4}-R_C\text{-heteroaryl}-R_{101}-R_1\text{-heterocycle}$,
 5 (XI) - $(CR_{C-x}R_{C-y})_{0-4}-R_1\text{-heterocycle}-R_{101}-R_C\text{-aryl}$,
 (XII) - $(CR_{C-x}R_{C-y})_{0-4}-R_1\text{-heterocycle}-R_{101}-R_C\text{-heteroaryl}$,
 (XIII) - $(CR_{C-x}R_{C-y})_{0-4}-R_1\text{-heterocycle}-R_{101}-R_1\text{-heterocycle}$, wherein
 R_{101} is a bond, $(CH_2)_{0-4}$, $-O-$, $-NH-$, or $-N(C_1-C_6 \text{ alkyl})$
 (XIV) - $(CR_{C-x}R_{C-y})_{0-4}-R_1\text{-heterocycle}$,
 10 (XV) - $[C(R_{C-1})(R_{C-2})]_{1-3}-CO-N(R_{C-3})_2$ where R_{C-1} and R_{C-2} are the
 same or different and are selected from the group consisting of:
 (A) $-H$,
 (B) $-C_1-C_6$ alkyl, optionally substituted with one, two
 or three substituents independently selected from the group
 15 consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$,
 $-CF_3$, C_1-C_6 alkoxy, $-O\text{-phenyl}$, and $-NR_{1-a}R_{1-b}$,
 (C) C_2-C_6 alkenyl with one or two double bonds,
 optionally substituted with one, two or three substituents
 independently selected from the group consisting of C_1-C_3 alkyl,
 20 $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O\text{-phenyl}$,
 and $-NR_{1-a}R_{1-b}$,
 (D) C_2-C_6 alkynyl optionally substituted with one, two
 or three substituents independently selected from the group
 consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$,
 25 $-CF_3$, C_1-C_6 alkoxy, $-O\text{-phenyl}$, and $-NR_{1-a}R_{1-b}$,
 (E) $-(CH_2)_{1-2}-S(O)_{0-2}-(C_1-C_6 \text{ alkyl})$,
 (F) $-(CH_2)_{0-4}-C_3-C_7$ cycloalkyl, optionally substituted
 with one, two or three substituents independently selected from
 the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$,
 30 $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O\text{-phenyl}$, and $-NR_{1-a}R_{1-b}$,
 (G) $-(C_1-C_4 \text{ alkyl})-R_1\text{-aryl}$,
 (H) $-(C_1-C_4 \text{ alkyl})-R_C\text{-heteroaryl}$,
 (I) $-(C_1-C_4 \text{ alkyl})-R_1\text{-heterocycle}$,
 (J) $-R_C\text{-heteroaryl}$,

(K) $-R_1\text{-heterocycle}$,

(M) $-(CH_2)_{1-4}-R_{C-4}-(CH_2)_{0-4}-R_1\text{-aryl}$ where R_{C-4} is $-O-$, $-S-$
or

$-NR(C_1-C_6 \text{ alkyl})-$,

5 (N) $-(CH_2)_{1-4}-R_{C-4}-(CH_2)_{0-4}-R_{C\text{-heteroaryl}}$,

(O) $-R_1\text{-aryl}$,

and where

R_{C-3} at each occurrence is independently:

(A) $-H$,

10 (B) $-C_1-C_6$ alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O\text{-phenyl}$, and $-NR_{1-a}R_{1-b}$,

(C) C_2-C_6 alkenyl with one or two double bonds,
15 optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, halogen, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O\text{-phenyl}$, and $-NR_{1-a}R_{1-b}$,

(D) C_2-C_6 alkynyl with one or two triple bonds,
20 optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O\text{-phenyl}$, and $-NR_{1-a}R_{1-b}$,

(E) $-(CH_2)_{0-4}-C_3-C_7$ cycloalkyl, optionally
25 substituted with 1, 2, or 3 groups that are independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O\text{-phenyl}$, and $-NR_{1-a}R_{1-b}$,

(F) $-R_1\text{-aryl}$,

30 (G) $-R_{C\text{-heteroaryl}}$,

(H) $-R_1\text{-heterocycle}$,

(I) $-(C_1-C_4 \text{ alkyl})-R_1\text{-aryl}$,

(J) $-(C_1-C_4 \text{ alkyl})-R_{C\text{-heteroaryl}}$,

(K) $-(C_1-C_4 \text{ alkyl})-R_1\text{-heterocycle}$,

(XVI) $-\text{CH}(\text{R}_{\text{C-aryl}})_2$,

(XVII) $-\text{CH}(\text{R}_{\text{C-heteroaryl}})_2$,

(XVIII) $-\text{CH}(\text{R}_{\text{C-aryl}})(\text{R}_{\text{C-heteroaryl}})$,

(XIX) -cyclopentyl, -cyclohexyl, or -cycloheptyl ring
 5 fused to $\text{R}_{\text{C-aryl}}$ or $\text{R}_{\text{C-heteroaryl}}$ or $\text{R}_{\text{1-heterocycle}}$ where $\text{R}_{\text{C-aryl}}$ or $\text{R}_{\text{C-heteroaryl}}$ or $\text{R}_{\text{1-heterocycle}}$ are as defined above where one carbon of cyclopentyl, cyclohexyl, or -cycloheptyl is optionally replaced with NH, $\text{NR}_{\text{N-5}}$, O, $\text{S}(=\text{O})_{0-2}$, and where cyclopentyl, cyclohexyl, or -cycloheptyl can be optionally substituted with one or two
 10 $-\text{C}_1-\text{C}_3$ alkyl, -F, -OH, -SH, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_6 alkoxy, =O, or $-\text{NR}_{1-a}\text{R}_{1-b}$,

(XX) C_2-C_{10} alkenyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-\text{C}\equiv\text{N}$,
 15 $-\text{CF}_3$, C_1-C_6 alkoxy, -O-phenyl, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(XXI) C_2-C_{10} alkynyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-\text{C}\equiv\text{N}$,
 $-\text{CF}_3$, C_1-C_6 alkoxy, -O-phenyl, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

20 (XXI) $-(\text{CH}_2)_{0-1}-\text{CHR}_{\text{C-6}}-(\text{CH}_2)_{0-1}-\text{R}_{\text{C-aryl}}$ wherein

$\text{R}_{\text{C-6}}$ is $-(\text{CH}_2)_{0-6}-\text{OH}$,

(XXII) $-(\text{CH}_2)_{0-1}-\text{CHR}_{\text{C-6}}-(\text{CH}_2)_{0-1}-\text{R}_{\text{C-heteroaryl}}$,

(XXIII) $-\text{CH}(-\text{R}_{\text{C-aryl}}$ or $\text{R}_{\text{C-heteroaryl}})-\text{CO}-\text{O}(\text{C}_1-\text{C}_4 \text{ alkyl})$,

(XXIV) $-\text{CH}(-\text{CH}_2-\text{OH})-\text{CH}(-\text{OH})-\text{alkyl}-\text{NO}_2$,

25 (XXV) $-(\text{C}_1-\text{C}_6 \text{ alkyl})-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})-\text{OH}$,

(XXVII) $-\text{CH}_2-\text{NH}-\text{CH}_2-\text{CH}(-\text{O}-\text{CH}_2-\text{CH}_3)_2$,

(XXVIII) -H, and

(XXIX) $-(\text{CH}_2)_{0-6}-\text{C}(=\text{NR}_{1-a})(\text{NR}_{1-a}\text{R}_{1-b})$;

where R_{N} is

30 (I) $\text{R}_{\text{N-1}}-\text{X}_{\text{N}}-$ where X_{N} is selected from the group consisting of:

(A) $-\text{CO}-$,

(B) $-\text{SO}_2-$,

(C) $-(\text{CR}'\text{'R}'\text{'})_{1-6}$ wherein

R''' and R''' at each occurrence are the same or different and are -H or C₁-C₄ alkyl,

(D) -CO-(CR''R''')₁₋₆-X_{N-1} wherein

X_{N-1} is selected from the group consisting of -O-,
5 -S- and -NR''-,

(E) a single bond, and

(F) -CO-(CR''R''')₁₋₆-

where R_{N-1} is selected from the group consisting of:

(A) R_{N-aryl} wherein R_{N-aryl} at each occurrence is
10 independently phenyl; naphthyl; tetralinyl; indanyl; indenyl;
dihydronaphthyl; or 6,7,8,9-tetrahydro-5H-benzo[a]cycloheptenyl;
each of which is optionally substituted with one, two or three
of the following substituents which can be the same or different
and are:

15 (1) C₁-C₆ alkyl, optionally substituted with one,
two or three substituents selected from the group consisting of
C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃
alkoxy, and -NR_{1-a}R_{1-b},

wherein R_{1-a} and R_{1-b} at each occurrence are
20 independently H or C₁-C₆ alkyl,

(2) -OH,

(3) -NO₂,

(4) -F, -Cl, -Br, -I,

(5) -CO₂H,

25 (6) -C≡N,

(7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are the
same or different and are selected from the group consisting of:

(a) -H,

(b) -C₁-C₈ alkyl optionally substituted with
30 one substituent selected from the group consisting of:

(i) -OH,

(ii) -NH₂,

(iii) phenyl,

(c) -C₁-C₈ alkyl optionally substituted with
35 1, 2, or 3 groups that are independently -F, -Cl, -Br, or -I,

(d) $-C_3-C_8$ cycloalkyl,
(e) $-(C_1-C_2 \text{ alkyl})-(C_3-C_8 \text{ cycloalkyl})$,
(f) $-(C_1-C_6 \text{ alkyl})-O-(C_1-C_3 \text{ alkyl})$,
(g) $-C_2-C_6$ alkenyl,
5 (h) $-C_2-C_6$ alkynyl,
(i) $-C_1-C_6$ alkyl chain with one double bond
and one triple bond,

(j) $-R_1\text{-aryl}$, wherein $R_1\text{-aryl}$ at each occurrence
is independently phenyl, naphthyl, indanyl, indenyl,
10 dihydronaphthyl, or tetralinyl each of which is optionally
substituted with 1, 2, 3, or 4 groups that are independently:

(i) C_1-C_6 alkyl optionally
substituted with one, two or three substituents independently
selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$,
15 $-I$, $-OH$, $-SH$, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

(ii) C_2-C_6 alkenyl with one or two
double bonds, optionally substituted with one, two or three
substituents independently selected from the group consisting of
 $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

20 (iii) C_2-C_6 alkynyl optionally
substituted with 1, 2, or 3 groups that are independently
selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(iv) $-F$, Cl , $-Br$ and $-I$,

25 (v) $-C_1-C_6$ alkoxy optionally
substituted with 1, 2, or 3 $-F$,

(vi) $-NR_{N-2}R_{N-3}$,

(vii) $-OH$,

(viii) $-C\equiv N$,

30 (ix) C_3-C_7 cycloalkyl, optionally
substituted with 1, 2, or 3 groups that are selected from the
group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy,
and $-NR_{1-a}R_{1-b}$,

(x) $-CO-(C_1-C_4 \text{ alkyl})$,

(xi) $-\text{SO}_2-\text{NR}_{1-a}\text{R}_{1-b}$,

(xii) $-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$, or

(xiii) $-\text{SO}_2-(\text{C}_1-\text{C}_4 \text{ alkyl})$,

(k) $-\text{R}_{1-\text{heteroaryl}}$, wherein $\text{R}_{1-\text{heteroaryl}}$ at each
5 occurrence is independently selected from the group consisting
of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl,
indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl,
quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl,
isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, 10
indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl,
furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl,
triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl,
isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-
carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl,
15 isoindolinyl, isobenzotetrahydrofuranyl,
isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl,
pyridopyridinyl, benzotetrahydrofuranyl, benzotetrahydrothienyl,
purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl,
pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
20 dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl,
coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-
oxide, tetrahydroquinolinyl, dihydroquinolinyl,
dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
25 dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl,
benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,
indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide,
quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-
30 oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
thiazolyl N-oxide, indoliziny N-oxide, indazolyl N-oxide,
benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and
35 benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(i) C_1 - C_6 alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C_1 - C_3 alkoxy,

(ii) C_2 - C_6 alkenyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C_1 - C_3 alkoxy, or -NR_{1-a}R_{1-b},

(iii) C_2 - C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C_1 - C_3 alkoxy, or -NR_{1-a}R_{1-b},

(iv) -F, -Cl, -Br and -I,

(v) C_1 - C_6 alkoxy optionally substituted with one, two, or three -F,

(vi) -(CH₂)₀₋₄-NR_{N-2}R_{N-3},

(vii) -OH,

(viii) -C≡N,

(ix) (CH₂)₀₋₄-C₃-C₇ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

(x) (CH₂)₀₋₄-CO-(C_1 - C_6 alkyl),

(xi) (CH₂)₀₋₄-SO₂-NR_{N-2}R_{N-3},

(xii) (CH₂)₀₋₄-CO-NR_{N-2}R_{N-3},

(xiii) (CH₂)₀₋₄-SO₂-(C_1 - C_6 alkyl),

(xiv) (CH₂)₀₋₄-N(R_{N-2})-SO₂-, and

(xv) (CH₂)₀₋₄-N(R_{N-2})-C(O)-,

(1) -R₁-heterocycle, wherein

R₁-heterocycle at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl,

tetrahydrothienyl, homopiperidinyl, homomorpholinyl,
homothiomorpholinyl, homothiomorpholinyl S,S-dioxide,
oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl,
dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl,
5 dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide,
tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the R_1 -heterocycle group is bonded by
any atom of the parent R_1 -heterocycle group substituted by hydrogen
such that the new bond to the R_1 -heterocycle group replaces the
10 hydrogen atom and its bond, where heterocycle is optionally
substituted with 1, 2, 3, or 4 groups that are independently:

(a) C_1-C_6 alkyl optionally
substituted with one, two or three substituents independently
selected from the group consisting of C_1-C_3 alkyl, halogen, -OH,
15 -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

(b) C_2-C_6 alkenyl with one or two
double bonds, optionally substituted with one, two or three
substituents independently selected from the group consisting of
-F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

20 (c) C_2-C_6 alkynyl with one or two
triple bonds, optionally substituted with one, two or three
substituents independently selected from the group consisting of
-F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(d) halogen,

25 (e) C_1-C_6 alkoxy,

(f) $-C_1-C_6$ alkoxy optionally
substituted with one, two, or three -F,

(g) $-NR_{N-2}R_{N-3}$,

(h) -OH,

30 (i) $-C\equiv N$,

(j) $(CH_2)_{0-4}-(C_3-C_8$ cycloalkyl),
optionally substituted with 1, 2, or 3 groups independently
selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, -
 CF_3 , C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

5

- (k) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_4 \text{ alkyl}),$
- (l) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{1-a}\text{R}_{1-b},$
- (m) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b},$
- (n) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_6 \text{ alkyl}),$ and
- (o) $=\text{O},$
- (p) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{N-2})-\text{SO}_2-$
- (q) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{N-2})-\text{C}(\text{O})-$

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- (8) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_{12} \text{ alkyl}),$
- (9) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkenyl}),$
- (10) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkynyl}),$
- (11) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl}),$
- (12) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-aryl},$
- (13) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl},$
- (14) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle},$

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(15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{N-4}$ wherein R_{N-4} is selected from the group consisting of phenyl, morpholinyl, thiomorpholinyl, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl, thienyl, pyrazolyl, pyridyl N-oxide, oxazolyl, thiazolyl, imidazolyl, and pyrrolidinyl where each group is optionally substituted with one, two, three, or four groups that are independently C_1-C_6 alkyl,

20

(16) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{N-5}$ where R_{N-5} is selected from the group consisting of:

25

- (a) $\text{C}_1-\text{C}_6 \text{ alkyl},$
- (b) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-aryl}),$
- (c) $\text{C}_2-\text{C}_6 \text{ alkenyl},$
- (d) $\text{C}_2-\text{C}_6 \text{ alkynyl},$
- (e) $-(\text{CH}_2)_{0-2}-\text{C}_3-\text{C}_8 \text{ cycloalkyl},$
- (f) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heteroaryl}),$ and
- (g) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heterocycle}),$

30

- (17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{N-2}\text{R}_{N-3},$
- (18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl}),$
- (19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl}),$
- (20) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3-\text{C}_8 \text{ cycloalkyl}),$

35

- (21) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{O}-\text{R}_{\text{N}-5},$
 (22) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2,$
 (23) $-(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2,$
 (24) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2},$
 5 (25) $-(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3},$
 (26) $-(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4},$
 (27) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl}),$
 (28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$ wherein
 R_{100} at each occurrence is independently -H
 10 or C_1-C_4 alkyl,
 (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2,$
 (30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2,$
 (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5}),$
 (32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH},$
 15 (33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5}),$
 (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl optionally}$
 substituted with one, two, three, four, or five of -F),
 (35) C_3-C_8 cycloalkyl,
 (36) C_2-C_6 alkenyl optionally substituted with C_1-
 20 C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy,
 or $-\text{NR}_{1-a}\text{R}_{1-b},$
 (37) C_2-C_6 alkynyl optionally substituted with C_1-
 C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy,
 or $-\text{NR}_{1-a}\text{R}_{1-b},$
 25 (38) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2},$
 (39) $-(\text{CH}_2)_{1-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl}),$
 (B) $-\text{R}_{\text{N-heteroaryl}}$ where $\text{R}_{\text{N-heteroaryl}}$ is selected from the
 group consisting of:
 pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl,
 30 indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl,
 quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl,
 isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indolizinyll,
 indazolyl, benzisothiazolyl, benzimidazolyl, benzofuranyl,
 furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl,
 35 triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl,

isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanlyl, chromanlyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, henoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazothiazolyl, dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl, dihydrobenzisothiazinyl, benzopyranlyl, benzothiopyranlyl, coumarinyl, isocoumarinyl, chromonyl, chromanonyl, tetrahydroquinolinyl, dihydroquinolinyl, dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl, dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl, benzoxazolinonyl, pyridinyl-N-oxide, pyrrolyl N-oxide, pyrimidinyl N-oxide, pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide, indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide, quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide, thiazolyl N-oxide, indolizinyl N-oxide, indazolyl N-oxide, benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-oxide, tetrazolyl N-oxide, benzothiopyranlyl S-oxide, benzothiopyranlyl S,S-dioxide, imidazopyrazolyl, quinazolinonyl, pyrazopyridyl, benzooxadiazolyl, dihydropyrimidinonyl, and dihydrobenzofuranonyl,

where the $R_{N\text{-heteroaryl}}$ group is bonded by any atom of the parent $R_{N\text{-heteroaryl}}$ group substituted by hydrogen such that the new bond to the $R_{N\text{-heteroaryl}}$ group replaces the hydrogen atom and its bond, where heteroaryl is optionally substituted with one, two, three, or four of:

(1) $C_1\text{-}C_6$ alkyl, optionally substituted with one, two or three substituents selected from the group consisting of $C_1\text{-}C_3$ alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-C\equiv N$, $-CF_3$, $C_1\text{-}C_3$ alkoxy, and $-NR_{1-a}R_{1-b}$,

(2) -OH,

- (3) $-\text{NO}_2$,
- (4) $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$,
- (5) $-\text{CO}_2\text{H}$,
- (6) $-\text{C}\equiv\text{N}$,
- 5 (7) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
- (8) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
- (9) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkenyl})$,
- (10) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkynyl})$,
- (11) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
- 10 (12) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-aryl}$,
- (13) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl}$,
- (14) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle}$,
- (15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$,
- (16) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$,
- 15 (17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
- (18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl})$,
- (19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
- (20) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
- (21) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$,
- 20 (22) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (23) $-(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (24) $-(\text{CH}_2)_{0-4}-\text{N}(-\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2}$,
- (25) $-(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
- (26) $-(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4}$,
- 25 (27) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
- (28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$,
- (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
- 30 (32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
- (33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
- (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})$ optionally substituted with one, two, three, four, or five of $-\text{F}$),
- (35) $\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,

(36) C₂-C₆ alkenyl optionally substituted with C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, or -NR_{1-a}R_{1-b},

(37) C₂-C₆ alkynyl optionally substituted with C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, or -NR_{1-a}R_{1-b},

(38) -(CH₂)₀₋₄-N(-H or R_{N-5})-SO₂-R_{N-2},

(39) -(CH₂)₁₋₄-C₃-C₈ cycloalkyl,

(C) R_{N-aryl}-W-R_{N-aryl},

(D) R_{N-aryl}-W-R_{N-heteroaryl},

(E) R_{N-aryl}-W-R_{1-heterocycle},

(F) R_{N-heteroaryl}-W-R_{N-aryl},

(G) R_{N-heteroaryl}-W-R_{N-heteroaryl},

(H) R_{N-heteroaryl}-W-R_{N-1-heterocycle},

(I) R_{N-heterocycle}-W-R_{N-aryl},

(J) R_{N-heterocycle}-W-R_{N-heteroaryl},

(K) R_{N-heterocycle}-W-R_{N-1-heterocycle},

where W is

(1) -(CH₂)₁₋₄-,

(2) -O-,

(3) -S(O)₀₋₂-,

(4) -N(R_{N-5})-,

(5) -CO-; or

(6) a bond;

(II) -CO-(C₁-C₁₀ alkyl) wherein the alkyl is optionally substituted with one two or three substituents independently selected from the group consisting of:

(A) -OH,

(B) -C₁-C₆ alkoxy,

(C) -C₁-C₆ thioalkoxy,

(D) -CO-O-R_{N-8} where

R_{N-8} at each occurrence is independently -H, C₁-C₆ alkyl or -phenyl,

(E) -CO-NR_{N-2}R_{N-3},

(F) -CO-R_{N-4},

- (G) $-\text{SO}_2-(\text{C}_1-\text{C}_8 \text{ alkyl})$,
(H) $-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
(I) $-\text{NH}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
(J) $-\text{NH}-\text{CO}-\text{O}-\text{R}_{\text{N}-8}$,
5 (K) $-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
(L) $-\text{R}_{\text{N}-4}$,
(M) $-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
(N) $-\text{O}-\text{CO}-\text{NR}_{\text{N}-8}\text{R}_{\text{N}-8}$,
(O) $-\text{O}-(\text{C}_1-\text{C}_5 \text{ alkyl})-\text{COOH}$,
10 (P) $-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})$ optionally substituted with one,
two, or three of $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$),
(Q) $-\text{NH}-\text{SO}_2-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
(R) halogen,
(S) $-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$,
15 (T) $-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-(\text{R}_{\text{N}-2})$, and
(U) $-\text{SO}_2-\text{R}_{\text{N}-2}$,

(III) $-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})$ wherein each alkyl is unsubstituted or independently substituted with one, two, or three substituents selected from the group consisting of :

- 20 (A) $-\text{OH}$,
(B) $-\text{C}_1-\text{C}_6 \text{ alkoxy}$,
(C) $-\text{C}_1-\text{C}_6 \text{ thioalkoxy}$,
(D) $-\text{CO}-\text{O}-\text{R}_{\text{N}-8}$,
(E) $-\text{CO}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
25 (F) $-\text{CO}-\text{R}_{\text{N}-4}$,
(G) $-\text{SO}_2-(\text{C}_1-\text{C}_8 \text{ alkyl})$,
(H) $-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
(I) $-\text{NH}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
(J) $-\text{NH}-\text{CO}-\text{O}-\text{R}_{\text{N}-8}$,
30 (K) $-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
(L) $-\text{R}_{\text{N}-4}$,
(M) $-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
(N) $-\text{O}-\text{CO}-\text{NR}_{\text{N}-8}\text{R}_{\text{N}-8}$,
(O) $-\text{O}-(\text{C}_1-\text{C}_5 \text{ alkyl})-\text{CO}_2\text{H}$,

(P) -O-(C₁-C₆ alkyl optionally substituted with one, two, or three groups that are independently -F, -Cl, -Br, or -I),

(Q) -NH-SO₂-(C₁-C₆ alkyl),

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(R) halogen,

(S) -N(H or R_{N-5})-SO₂-R_{N-2},

(T) -N(H or R_{N-5})-CO-(R_{N-2}), and

(U) -SO₂-R_{N-2},

(IV) -CO-(C₁-C₆ alkyl)-S-(C₁-C₆ alkyl) wherein each
10 alkyl is unsubstituted or substituted with one, two, or three of
substituents independently selected from the group consisting
of:

(A) -OH,

(B) -C₁-C₆ alkoxy,

15

(C) -C₁-C₆ thioalkoxy,

(D) -CO-O-R_{N-8},

(E) -CO-NR_{N-2}R_{N-3},

(F) -CO-R_{N-4},

(G) -SO₂-(C₁-C₈ alkyl),

20

(H) -SO₂-NR_{N-2}R_{N-3},

(I) -NH-CO-(C₁-C₆ alkyl),

(J) -NH-CO-O-R_{N-8},

(K) -NR_{N-2}R_{N-3},

(L) -R_{N-4},

25

(M) -O-CO-(C₁-C₆ alkyl),

(N) -O-CO-NR_{N-8}R_{N-8},

(O) -O-(C₁-C₅ alkyl)-COOH,

(P) -O-(C₁-C₆ alkyl optionally substituted with
one, two, or three groups that are independently -F, -Cl, -Br,
30 or -I),

(Q) -NH-SO₂-(C₁-C₆ alkyl),

(R) halogen,

(S) -N(H or R_{N-5})-SO₂-R_{N-2},

(T) -N(H or R_{N-5})-CO-(R_{N-2}), and

(U) $-\text{SO}_2-\text{R}_{\text{N}-2}$,(V) $-\text{CO}-\text{CH}(-(\text{CH}_2)_{0-2}-\text{O}-\text{R}_{\text{N}-10})-(\text{CH}_2)_{0-2}-\text{R}_{\text{N-aryl}}/\text{R}_{\text{N-heteroaryl}}$

wherein

 $\text{R}_{\text{N}-10}$ is selected from the group consisting of:

5

(A) $-\text{H}$,(B) C_1-C_6 alkyl,(C) C_3-C_8 cycloalkyl,(D) C_2-C_6 alkenyl with one double bond,(E) C_2-C_6 alkynyl with one triple bond,

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(F) $\text{R}_1\text{-aryl}$,(G) $\text{R}_{\text{N-heteroaryl}}$,(H) $\text{R}_{\text{N-heterocycle}}$,

(VI) $-\text{CO}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$ where the cycloalkyl group is optionally substituted with one or two substituents independently selected from the group consisting of:

15

(A) $-(\text{CH}_2)_{0-4}-\text{OH}$,(B) $-(\text{CH}_2)_{0-4}-\text{C}_1-\text{C}_6$ alkoxy,(C) $-(\text{CH}_2)_{0-4}-\text{C}_1-\text{C}_6$ thioalkoxy,(D) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{\text{N}-8}$,

20

(E) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,(F) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$,(G) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_8 \text{ alkyl})$,(H) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,(I) $-(\text{CH}_2)_{0-4}-\text{NH}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,

25

(J) $-\text{NH}-\text{CO}-\text{O}-\text{R}_{\text{N}-8}$,(K) $-(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,(L) $-(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4}$,(M) $-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,(N) $-\text{O}-\text{CO}-\text{NR}_{\text{N}-8}\text{R}_{\text{N}-8}$,

30

(O) $-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})-\text{CO}_2\text{H}$,

(P) $-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})$ optionally substituted with one, two, or three groups that are independently selected from $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, and $-\text{I}$,

- (Q) $-\text{NH}-\text{SO}_2-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
 (R) halogen,
 (S) $-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$, and
 (T) $-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-(\text{R}_{\text{N}-2})$, and
 5 (U) $-\text{SO}_2-\text{R}_{\text{N}-2}$;

where R_1 and R_2 are independently H, aryl, heteroaryl, or

(I) C_1-C_6 alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C_1-C_3 alkyl, C_1-C_7 alkyl (optionally substituted with C_1-C_3 alkyl and C_1-C_3 alkoxy), $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, $-\text{NR}_{1-a}\text{R}_{1-b}$ where R_{1-a} and R_{1-b} are independently $-\text{H}$ or C_1-C_6 alkyl, and $-\text{OC}=\text{O NR}_{1-a}\text{R}_{1-b}$,

(II) $-\text{CH}_2-\text{S}(\text{O})_{0-2}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,

(III) $-\text{CH}_2-\text{CH}_2-\text{S}(\text{O})_{0-2}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,

15 (IV) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(V) C_2-C_6 alkynyl with one or two triple bonds, 20 optionally substituted with one, two or three substituents selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(VI) $-(\text{CH}_2)_{n1}-(\text{R}_{1-\text{aryl}})$ where n_1 is zero or one and where $\text{R}_{1-\text{aryl}}$ is phenyl, 1-naphthyl, 2-naphthyl and indanyl, indenyl, 25 dihydronaphthalyl, or tetralinyl optionally substituted with one, two, three or four of the following substituents on the aryl ring:

(A) C_1-C_6 alkyl optionally substituted with one, two or three substituents selected from the group consisting of 30 C_1-C_3 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(B) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents

selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(C) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(D) -F, Cl, -Br or -I,

(F) -C₁-C₆ alkoxy optionally substituted with one, two or three of -F,

(G) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(H) -OH,

(I) -C≡N,

(J) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(K) -CO-(C₁-C₄ alkyl),

(L) -SO₂-NR_{1-a}R_{1-b},

(M) -CO-NR_{1-a}R_{1-b},

(N) -SO₂-(C₁-C₄ alkyl),

(VII) -(CH₂)_{n1}-(R_{1-heteroaryl}) where n₁ is as defined above and where R_{1-heteroaryl} is selected from the group consisting of:

pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indolizinyl, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl,

pyridopyridinyl, benzotetrahydrofuranyl, benzotetrahydrothienyl,
 purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl,
 pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
 dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
 5 dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl,
 coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-
 oxide, tetrahydroquinolinyl, dihydroquinolinyl,
 dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
 dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl,
 10 benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
 pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,
 indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide,
 quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-
 oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
 15 thiazolyl N-oxide, indolizinyl N-oxide, indazolyl N-oxide,
 benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
 oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
 oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide,
 benzothiopyranyl S,S-dioxide,

20 where the $R_{1\text{-heteroaryl}}$ group is bonded to $-(CH_2)_{n1}-$ by any ring atom
 of the parent $R_{N\text{-heteroaryl}}$ group substituted by hydrogen such that
 the new bond to the $R_{1\text{-heteroaryl}}$ group replaces the hydrogen atom
 and its bond, where heteroaryl is optionally substituted with
 one, two, three or four of:

25 (1) C_1-C_6 alkyl optionally substituted with one,
 two or three substituents selected from the group consisting of
 C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH,
 -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(2) C_2-C_6 alkenyl with one or two double bonds,
 30 optionally substituted with one, two or three substituents
 selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, -
 CF_3 , C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(3) C_2-C_6 alkynyl with one or two triple bonds,
 optionally substituted with one, two or three substituents

selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(4) -F, Cl, -Br or -I,

(6) -C₁-C₆ alkoxy optionally substituted with
5 one, two, or three of -F,

(7) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined
below,

(8) -OH,

(9) -C≡N,

(10) C₃-C₇ cycloalkyl, optionally substituted
10 with one, two or three substituents selected from the group
consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -
NR_{1-a}R_{1-b},

(11) -CO-(C₁-C₄ alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

(13) -CO-NR_{1-a}R_{1-b}, or

(14) -SO₂-(C₁-C₄ alkyl), with the proviso that
15 when n₁ is zero R_{1-heteroaryl} is not bonded to the carbon chain by
nitrogen, or

(VIII) -(CH₂)_{n1}-(R_{1-heterocycle}) where n₁ is as defined
20 above and R_{1-heterocycle} is selected from the group consisting of:

morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide,
thiomorpholinyl S,S-dioxide, piperazinyl,
homopiperazinyl, pyrrolidinyl, pyrrolinyl,
25 tetrahydropyranyl, piperidinyl, tetrahydrofuranyl,
tetrahydrothienyl, homopiperidinyl, homomorpholinyl,
homomorpholinyl S-oxide, homothiomorpholinyl S,S-
dioxide, oxazolidinonyl, dihydropyrazolyl,
dihydropyrrolyl dihydropyrazinyl dihydropyridinyl
30 dihydropyrimidinyl, dihydrofuryl, dihydropyranyl,
tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-
dioxide, homothiomorpholinyl S-oxide,

where the R_{1-heterocycle} group is bonded by any atom of the parent
R_{1-heterocycle} group substituted by hydrogen such that the new bond

to the R_1 -heterocycle group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with one, two, three or four:

5 (1) C_1 - C_6 alkyl optionally substituted with one, two or three substituents selected from the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

10 (2) C_2 - C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

15 (3) C_2 - C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

(4) -F, Cl, -Br, or -I,

(5) C_1 - C_6 alkoxy,

(6) - C_1 - C_6 alkoxy optionally substituted with one, two, or three -F,

20 (7) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(8) -OH,

(9) -C \equiv N,

25 (10) C_3 - C_7 cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

(11) -CO-(C_1 - C_4 alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

30 (13) -CO-NR_{1-a}R_{1-b},

(14) -SO₂-(C_1 - C_4 alkyl),

(15) =O, with the proviso that when n_1 is zero R_1 -heterocycle is not bonded to the carbon chain by nitrogen; and

where R_{20} is H or C_{1-6} alkyl or alkenyl.

In alternative broad aspect, the invention provides compounds of formula I, wherein each of the R_2 groups (I)-(VIII) is attached to the nitrogen carrying R_{20} by -Z-, wherein Z is -
5 C(O)-, -CO₂ or -SO₂-.

The invention also provides compounds, compositions, kits, and methods for inhibiting beta-secretase-mediated cleavage of amyloid precursor protein (APP). More particularly, the compounds, compositions, and methods of the invention are
10 effective to inhibit the production of A-beta peptide and to treat and/or prevent any human or veterinary disease or condition associated with a pathological form of A-beta peptide.

The invention also include methods of treating a patient who has, or in preventing a patient from getting, a disease or
15 condition selected from the group consisting of Alzheimer's disease, for helping prevent or delay the onset of Alzheimer's disease, for treating patients with mild cognitive impairment (MCI) and preventing or delaying the onset of Alzheimer's disease in those who would progress from MCI to AD, for treating
20 Down's syndrome, for treating humans who have Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, for treating cerebral amyloid angiopathy and preventing its potential consequences, i.e. single and recurrent lobar hemorrhages, for treating other degenerative dementias,
25 including dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's disease, frontotemporal dementias with parkinsonism (FTDP), dementia associated with progressive supranuclear palsy, dementia associated with cortical basal degeneration, or diffuse Lewy body type of
30 Alzheimer's disease and who is in need of such treatment, which includes administration of a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof.

The compounds of the invention possess beta-secretase
35 inhibitory activity. The inhibitory activities of the compounds

of the invention is readily demonstrated, for example, using one or more of the assays described herein or known in the art.

In an embodiment, this method of treatment can be used where the disease is Alzheimer's disease.

5 In an embodiment, this method of treatment can help prevent or delay the onset of Alzheimer's disease.

In an embodiment, this method of treatment can be used where the disease is mild cognitive impairment.

10 In an embodiment, this method of treatment can be used where the disease is Down's syndrome.

In an embodiment, this method of treatment can be used where the disease is Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type.

15 In an embodiment, this method of treatment can be used where the disease is cerebral amyloid angiopathy.

In an embodiment, this method of treatment can be used where the disease is degenerative dementias.

20 In an embodiment, this method of treatment can be used where the disease is diffuse Lewy body type of Alzheimer's disease.

In an embodiment, this method of treatment can treat an existing disease.

In an embodiment, this method of treatment can prevent a disease from developing.

25 In an embodiment, this method of treatment can employ therapeutically effective amounts: for oral administration from about 0.1 mg/day to about 1,000 mg/day; for parenteral, sublingual, intranasal, intrathecal administration from about 0.5 to about 100 mg/day; for depo administration and implants
30 from about 0.5 mg/day to about 50 mg/day; for topical administration from about 0.5 mg/day to about 200 mg/day; for rectal administration from about 0.5 mg to about 500 mg.

In an embodiment, this method of treatment can employ therapeutically effective amounts: for oral administration from

about 1 mg/day to about 100 mg/day; and for parenteral administration from about 5 to about 50 mg daily.

In an embodiment, this method of treatment can employ therapeutically effective amounts for oral administration from about 5 mg/day to about 50 mg/day.

The invention also includes a pharmaceutical composition which includes a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof.

The invention also includes the use of a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof, for the manufacture of a medicament for use in treating a patient who has, or in preventing a patient from getting, a disease or condition selected from the group consisting of Alzheimer's disease, for helping prevent or delay the onset of Alzheimer's disease, for treating patients with mild cognitive impairment (MCI) and preventing or delaying the onset of Alzheimer's disease in those who would progress from MCI to AD, for treating Down's syndrome, for treating humans who have Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, for treating cerebral amyloid angiopathy and preventing its potential consequences, i.e. single and recurrent lobar hemorrhages, for treating other degenerative dementias, including dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's disease, dementia associated with progressive supranuclear palsy, dementia associated with cortical basal degeneration, diffuse Lewy body type of Alzheimer's disease and who is in need of such treatment.

In an embodiment, this use of a compound of the formula hereinabove can be employed where the disease is Alzheimer's disease.

In an embodiment, this use of a compound of the formula hereinabove can help prevent or delay the onset of Alzheimer's disease.

In an embodiment, this use of a compound of the formula hereinabove can be employed where the disease is mild cognitive impairment.

5 In an embodiment, this use of a compound of the formula hereinabove can be employed where the disease is Down's syndrome.

In an embodiment, this use of a compound of the formula hereinabove can be employed where the disease is Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type.

10 In an embodiment, this use of a compound of the formula hereinabove can be employed where the disease is cerebral amyloid angiopathy.

In an embodiment, this use of a compound of the formula hereinabove can be employed where the disease is degenerative
15 dementias.

In an embodiment, this use of a compound of the formula hereinabove can be employed where the disease is diffuse Lewy body type of Alzheimer's disease.

In an embodiment, this use of a substituted amine employs a
20 pharmaceutically acceptable salt selected from the group consisting of salts of the following acids hydrochloric, hydrobromic, hydroiodic, nitric, sulfuric, phosphoric, citric, methanesulfonic, $\text{CH}_3-(\text{CH}_2)_n-\text{COOH}$ where n is 0 through 4, $\text{HOOC}-(\text{CH}_2)_n-\text{COOH}$ where n is as defined above, $\text{HOOC}-\text{CH}=\text{CH}-\text{COOH}$, and
25 phenyl-COOH.

The invention also includes methods for inhibiting beta-secretase activity, for inhibiting cleavage of amyloid precursor protein (APP), in a reaction mixture, at a site between Met596 and Asp597, numbered for the APP-695 amino acid isotype, or at a
30 corresponding site of an isotype or mutant thereof; for inhibiting production of amyloid beta peptide (A beta) in a cell; for inhibiting the production of beta-amyloid plaque in an animal; and for treating or preventing a disease characterized by beta-amyloid deposits in the brain. These methods each
35 include administration of a therapeutically effective amount of

a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof.

The invention also includes a method for inhibiting beta-secretase activity, including exposing said beta-secretase to an effective inhibitory amount of a compound of formula, or a
5 pharmaceutically acceptable salt or ester thereof.

In an embodiment, this method employs a compound that inhibits 50% of the enzyme's activity at a concentration of less than 50 micromolar.

10 In an embodiment, this method employs a compound that inhibits 50% of the enzyme's activity at a concentration of 10 micromolar or less.

In an embodiment, this method employs a compound that inhibits 50% of the enzyme's activity at a concentration of 1
15 micromolar or less.

In an embodiment, this method employs a compound that inhibits 50% of the enzyme's activity at a concentration of 10 nanomolar or less.

In an embodiment, this method includes exposing said beta-secretase to said compound *in vitro*.
20

In an embodiment, this method includes exposing said beta-secretase to said compound in a cell.

In an embodiment, this method includes exposing said beta-secretase to said compound in a cell in an animal.

25 In an embodiment, this method includes exposing said beta-secretase to said compound in a human.

The invention also includes a method for inhibiting cleavage of amyloid precursor protein (APP), in a reaction mixture, at a site between Met596 and Asp597, numbered for the
30 APP-695 amino acid isotype; or at a corresponding site of an isotype or mutant thereof, including exposing said reaction mixture to an effective inhibitory amount of a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof.

In an embodiment, this method employs a cleavage site: between Met652 and Asp653, numbered for the APP-751 isotype; between Met 671 and Asp 672, numbered for the APP-770 isotype; between Leu596 and Asp597 of the APP-695 Swedish Mutation; 5 between Leu652 and Asp653 of the APP-751 Swedish Mutation; or between Leu671 and Asp672 of the APP-770 Swedish Mutation.

In an embodiment, this method exposes said reaction mixture *in vitro*.

10 In an embodiment, this method exposes said reaction mixture in a cell.

In an embodiment, this method exposes said reaction mixture in an animal cell.

In an embodiment, this method exposes said reaction mixture in a human cell.

15 The invention also includes a method for inhibiting production of amyloid beta peptide (A beta) in a cell, including administering to said cell an effective inhibitory amount of a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof.

20 In an embodiment, this method includes administering to an animal.

In an embodiment, this method includes administering to a human.

25 The invention also includes a method for inhibiting the production of beta-amyloid plaque in an animal, including administering to said animal an effective inhibitory amount of a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof.

30 In an embodiment, this method includes administering to a human.

The invention also includes a method for treating or preventing a disease characterized by beta-amyloid deposits in the brain including administering to a patient an effective therapeutic amount of a hydroxyethylene compound of formula (I), 35 or a pharmaceutically acceptable salt or ester thereof.

In an embodiment, this method employs a compound that inhibits 50% of the enzyme's activity at a concentration of less than 50 micromolar.

5 In an embodiment, this method employs a compound that inhibits 50% of the enzyme's activity at a concentration of 10 micromolar or less.

In an embodiment, this method employs a compound that inhibits 50% of the enzyme's activity at a concentration of 1 micromolar or less.

10 In an embodiment, this method employs a compound that inhibits 50% of the enzyme's activity at a concentration of 10 nanomolar or less.

In an embodiment, this method employs a compound at a therapeutic amount in the range of from about 0.1 to about 1000
15 mg/day.

In an embodiment, this method employs a compound at a therapeutic amount in the range of from about 15 to about 1500 mg/day.

In an embodiment, this method employs a compound at a
20 therapeutic amount in the range of from about 1 to about 100 mg/day.

In an embodiment, this method employs a compound at a therapeutic amount in the range of from about 5 to about 50 mg/day.

25 In an embodiment, this method can be used where said disease is Alzheimer's disease.

In an embodiment, this method can be used where said disease is Mild Cognitive Impairment, Down's Syndrome, or Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch
30 Type.

The invention also includes a composition including beta-secretase complexed with a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof.

The invention also includes a method for producing a beta-
35 secretase complex including exposing beta-secretase to a

compound of formula (I), or a pharmaceutically acceptable salt or ester thereof, in a reaction mixture under conditions suitable for the production of said complex.

In an embodiment, this method employs exposing *in vitro*.

5 In an embodiment, this method employs a reaction mixture that is a cell.

The invention also includes a component kit including component parts capable of being assembled, in which at least one component part includes a compound of formula I enclosed in
10 a container.

In an embodiment, this component kit includes lyophilized compound, and at least one further component part includes a diluent.

The invention also includes a container kit including a
15 plurality of containers, each container including one or more unit dose of a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof.

In an embodiment, this container kit includes each container adapted for oral delivery and includes a tablet, gel,
20 or capsule.

In an embodiment, this container kit includes each container adapted for parenteral delivery and includes a depot product, syringe, ampoule, or vial.

In an embodiment, this container kit includes each
25 container adapted for topical delivery and includes a patch, medipad, ointment, or cream.

The invention also includes an agent kit including a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof; and one or more therapeutic agent selected
30 from the group consisting of an antioxidant, an anti-inflammatory, a gamma secretase inhibitor, a neurotrophic agent, an acetyl cholinesterase inhibitor, a statin, an A beta peptide, and an anti-A beta antibody.

The invention also includes a composition including: a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof; and an inert diluent or edible carrier.

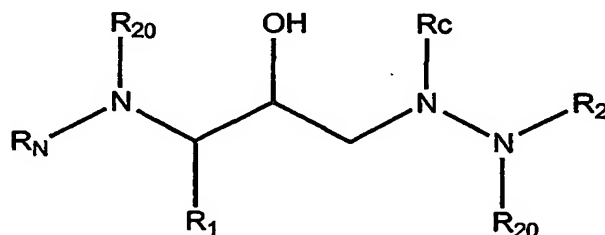
In an embodiment, this composition includes a carrier that is an oil.

The invention also includes a composition including: a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof; and a binder, excipient, disintegrating agent, lubricant, or gildant.

The invention also includes a composition including: a compound of formula (I), or a pharmaceutically acceptable salt or ester thereof; disposed in a cream, ointment, or patch.

Detailed Description of the Invention

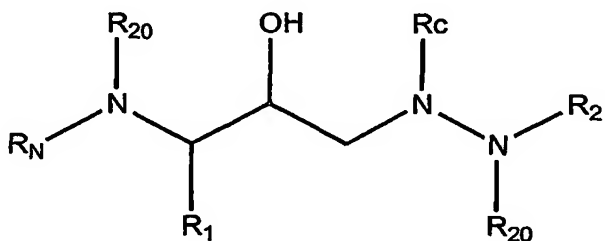
As noted above, the invention provides compounds of formula I:



(I)

where R_N, R_c, R₁, R₂ and R₂₀ are as defined above, and pharmaceutically acceptable salts and esters thereof.

In a preferred embodiment, the invention provides for compounds of formula II:



(II)

or a pharmaceutically acceptable salt thereof,
where R_c is

(I) $-C_1-C_{10}$ alkyl optionally substituted with one, two or three groups independently selected from the group consisting of
5 C_1-C_3 alkyl, halogen, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O$ -phenyl, $-NR_{1-a}R_{1-b}$, $-OC=O NR_{1-a}R_{1-b}$, $-S(=O)_{0-2} R_{1-a}$, $-NR_{1-a}C=O NR_{1-a}R_{1-b}$, $-C=O NR_{1-a}R_{1-b}$, and $-S(=O)_2 NR_{1-a}R_{1-b}$ wherein

R_{1-a} and R_{1-b} at each occurrence are independently H or C_1-C_6 alkyl,

10 (II) $-(CH_2)_{0-3}-(C_3-C_8)$ cycloalkyl where cycloalkyl can be optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, halogen, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_6 alkoxy, $-O$ -phenyl, $-CO_2H$, $-CO_2-(C_1-C_4)$ alkyl, and $-NR_{1-a}R_{1-b}$

15 (III) $-(CR_{c-x}R_{c-y})_{0-4}-R_{c-aryl}$ where R_{c-x} and R_{c-y} are independently selected from the group consisting of

$-H$,

C_1-C_4 alkyl optionally substituted with 1 or 2 $-OH$,

C_1-C_4 alkoxy optionally substituted with 1, 2, or 3

20 halogen,

$-(CH_2)_{0-4}-C_3-C_8$ cycloalkyl,

C_2-C_6 alkenyl containing one or two double bonds,

C_2-C_6 alkynyl containing one or two triple bonds, and
phenyl,

25 or

R_{c-x} and R_{c-y} are taken together with the carbon to which they are attached to form a carbocycle of three, four, five, six or seven carbon atoms, where one carbon atom is optionally replaced by a group selected from $-O-$, $-S-$, $-SO_2-$,
30 $-NR_{N-2}-$ and R_{c-aryl} , wherein

R_{c-aryl} is phenyl, which is optionally substituted with 1, 2, or 3 groups that are independently:

(1) C_1-C_6 alkyl, optionally substituted with one, two or three substituents selected from the group

consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(2) -OH,

(3) -NO₂,

5 (4) halogen,

(5) -CO₂H,

(6) -C≡N,

(7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3} where

R_{N-2} and R_{N-3} are independently selected

10 from the group consisting of:

(a) -H,

(b) -C₁-C₆ alkyl optionally substituted with one substituent selected from the group consisting of:

(i) -OH, and

15 (ii) -NH₂,

(c) -C₁-C₆ alkyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -Br, -I, or OH,

(d) -C₃-C₇ cycloalkyl,

20 (e) -(C₁-C₂ alkyl)-(C₃-C₇ cycloalkyl),

(f) -(C₁-C₆ alkyl)-O-(C₁-C₃ alkyl),

(g) -C₂-C₆ alkenyl

(h) -C₂-C₆ alkynyl

25 (i) -C₁-C₆ alkyl chain with one double bond and one triple bond,

(j) -R_{1-aryl} wherein R_{1-aryl} at each occurrence is independently phenyl, naphthyl, indanyl, indenyl, dihydronaphthyl, or tetralinyl each of which is optionally substituted with 1, 2, 3, or 4 groups that are independently:

30 (i) C₁-C₆ alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C₁-C₃ alkoxy,

(ii) C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

5 (iii) C₂-C₆ alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(iv) -F, Cl, -Br and -I,

10 (v) -C₁-C₆ alkoxy optionally substituted with 1, 2, or 3 -F,

(vi) -NR_{N-2}R_{N-3},

(vii) -OH,

(viii) -C≡N,

15 (ix) C₃-C₇ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(x) -CO-(C₁-C₄ alkyl),

20 (xi) -SO₂-NR_{1-a}R_{1-b},

(xii) -CO-NR_{1-a}R_{1-b}, or

(xiii) -SO₂-(C₁-C₄ alkyl),

(k) -R₁-heteroaryl wherein R₁-heteroaryl at each occurrence is independently selected from the group
25 consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indolizinyl, indazolyl, benzothiazolyl, benzimidazolyl,
30 benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl,

isobenzotetrahydrofuranyl, isobenzotetrahydrothienyl,
 isobenzothienyl, benzoxazolyl, pyridopyridinyl,
 benzotetrahydrofuranyl, benzotetrahydrothienyl, purinyl,
 benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl,
 5 pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
 dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
 dihydrobenzisoctiazinyl, benzopyranyl, benzothiopyranyl,
 coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-
 oxide, tetrahydroquinolinyl, dihydroquinolinyl,
 10 dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
 dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl,
 benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
 pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,
 indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide,
 15 quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-
 oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
 thiazolyl N-oxide, indolizinyl N-oxide, indazolyl N-oxide,
 benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
 oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
 20 oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and
 benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is optionally substituted with 1, 2, 3, or 4 groups that are independently:

- 25 (i) C_1 - C_6 alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C_1 - C_3 alkoxy,
- (ii) C_2 - C_6 alkenyl optionally
 30 substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},
- (iii) C_2 - C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are

independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(iv) -F, -Cl, -Br and -I,

5 substituted with one, two, or three -F,

(vi) -(CH₂)₀₋₄-NR_{N-2}R_{N-3},

(vii) -OH,

(viii) -C≡N,

10 optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(x) (CH₂)₀₋₄-CO-(C₁-C₆ alkyl),

(xi) (CH₂)₀₋₄-SO₂-NR_{N-2}R_{N-3},

15

(xii) (CH₂)₀₋₄-CO-NR_{N-2}R_{N-3},

(xiii) (CH₂)₀₋₄-SO₂-(C₁-C₆

alkyl),

(xiv) (CH₂)₀₋₄-N(R_{N-2})-SO₂-,

and

20

(xv) (CH₂)₀₋₄-N(R_{N-2})-C(O)-,

(8) -(CH₂)₀₋₄-CO-(C₁-C₁₂ alkyl),

(9) -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkenyl),

(10) -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkynyl),

(11) -(CH₂)₀₋₄-CO-(CH₂)₀₋₄ (C₃-C₇ cycloalkyl),

25

(12) -(CH₂)₀₋₄-CO-R_{1-aryl},

(13) -(CH₂)₀₋₄-CO-R_{1-heteroaryl},

(14) -(CH₂)₀₋₄-CO-R_{1-heterocycle} wherein

R_{1-heterocycle} at each occurrence is independently selected from the group consisting of morpholinyl, 30 thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S,S-dioxide,

oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl,
dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl,
dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide,
tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

5 where the R_1 -heterocycle group is bonded by
any atom of the parent R_1 -heterocycle group substituted by hydrogen
such that the new bond to the R_1 -heterocycle group replaces the
hydrogen atom and its bond, where heterocycle is optionally
substituted with 1, 2, 3, or 4 groups that are independently:

10 (a) C_1 - C_6 alkyl optionally
substituted with one, two or three substituents independently
selected from the group consisting of C_1 - C_3 alkyl, halogen, -OH,
-SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1 - C_3 alkoxy,

(b) C_2 - C_6 alkenyl with one or two
15 double bonds, optionally substituted with one, two or three
substituents independently selected from the group consisting of
-F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(c) C_2 - C_6 alkynyl with one or two
triple bonds, optionally substituted with one, two or three
20 substituents independently selected from the group consisting of
-F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(d) halogen,

(e) C_1 - C_6 alkoxy,

(f) $-C_1$ - C_6 alkoxy optionally
25 substituted with one, two, or three -F,

(g) $-NR_{N-2}R_{N-3}$,

(h) -OH,

(i) $-C\equiv N$,

(j) $(CH_2)_{0-4}$ -(C_3 - C_7 cycloalkyl),
30 optionally substituted with 1, 2, or 3 groups independently
selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, -
 CF_3 , C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(k) $-(CH_2)_{0-4}-CO-(C_1-C_4$ alkyl),

(l) $-(CH_2)_{0-4}-SO_2-NR_{1-a}R_{1-b}$,

- (m) $-(CH_2)_{0-4}-CO-NR_{1-a}R_{1-b}$,
 (n) $-(CH_2)_{0-4}-SO_2-(C_1-C_6 \text{ alkyl})$, and
 (o) $=O$,
 (p) $-(CH_2)_{0-4}-N(R_{N-2})-SO_2-$
 (q) $-(CH_2)_{0-4}-N(R_{N-2})-C(O)-$

5

(15) $-(CH_2)_{0-4}-CO-R_{N-4}$ wherein

R_{N-4} at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, pyrrolidinonyl, pyrrolyl, pyrazolyl, thienyl, pyridyl N-oxide, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl and pyrrolidinyl where each group is optionally substituted with 1, 2, 3, or 4 groups that are independently C_1-C_6 alkyl,

10

15

(16) $-(CH_2)_{0-4}-CO_2-R_{N-5}$ where

R_{N-5} at each occurrence is independently selected from the group consisting of:

- (a) C_1-C_6 alkyl,
 (b) $-(CH_2)_{0-2}-(R_{1-aryl})$,
 (c) C_2-C_6 alkenyl,
 (d) C_2-C_6 alkynyl,
 (e) C_3-C_7 cycloalkyl, and
 (f) $-(CH_2)_{0-4}-(R_{1-heteroaryl})$,

20

(17) $-(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$

25

(18) $-(CH_2)_{0-4}-SO-(C_1-C_8 \text{ alkyl})$,

(19) $-(CH_2)_{0-4}-SO_2-(C_1-C_{12} \text{ alkyl})$,

(20) $-(CH_2)_{0-4}-SO_2-(C_3-C_7 \text{ cycloalkyl})$,

(21) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO_2-R_{N-5}$,

(22) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO-N(R_{N-5})_2$,

30

(23) $-(CH_2)_{0-4}-N-CS-N(R_{N-5})_2$,

(24) $-(CH_2)_{0-4}-N(-H \text{ or } R_{N-5})-CO-R_{N-2}$,

(25) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,

(26) $-(CH_2)_{0-4}-R_{N-4}$,

(27) $-(CH_2)_{0-4}-O-CO-(C_1-C_6 \text{ alkyl})$,

(28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$ where R_{100} is independently H or C_1-C_4 alkyl,

(29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,

(30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,

5 (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,

(32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,

(33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,

10 (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})$ wherein the alkyl group is optionally substituted with one, two, three, four, or five substituents independently selected from the group consisting of F, Cl, Br, and I,

(35) $-(\text{CH}_2)_{0-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,

15 (36) C_2-C_6 alkenyl optionally substituted with C_1-C_3 alkyl, halogen, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, or $-\text{NR}_{1-a}\text{R}_{1-b}$,

(37) C_2-C_6 alkynyl optionally substituted with C_1-C_3 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, or $-\text{NR}_{1-a}\text{R}_{1-b}$, and

(38) $-(\text{CH}_2)_{0-4}-\text{N}(-\text{H} \text{ or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$;

20 (IV) $-(\text{CR}_{\text{C}-x}\text{R}_{\text{C}-y})_{0-4}-\text{R}_{\text{C-heteroaryl}}$ wherein $\text{R}_{\text{C-heteroaryl}}$ at each occurrence is independently selected from the group consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, 25 pyrazolyl, oxazolyl, thiazolyl, indolizinyll, indazolyl, benzoisothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, 30 tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, hexoxazinyl, phenothiazinyl, 35 pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,

dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl,
coumarinyl, isocoumarinyl, chromonyl, chromanonyl,
tetrahydroquinolinyl, dihydroquinolinyl,
5 dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
dihydroisocoumarinyl, isoindolinonyl,
benzodioxanyl, benzoxazolinonyl, imidazopyrazolyl,
quinazolinonyl, pyrazopyridyl, benzooxadiazolyl,
dihydropyrimidinonyl, dihydrobenzofuranonyl,

10 where the $R_{C\text{-heteroaryl}}$ group is bonded by any atom of the
parent $R_{C\text{-heteroaryl}}$ group substituted by hydrogen such that the new
bond to the $R_{C\text{-heteroaryl}}$ group replaces the hydrogen atom and its
bond, where heteroaryl is optionally substituted 1, 2, 3, or 4
groups that are independently:

15 (1) $C_1\text{-}C_6$ alkyl, optionally substituted with 1, 2, or
3 groups independently selected from the group consisting of $C_1\text{-}$
 C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF $_3$, $C_1\text{-}C_3$ alkoxy,
and -NR $_{1-a}$ R $_{1-b}$,

(2) -OH,

20 (3) -NO $_2$,

(4) -F, -Cl, -Br, -I,

(5) -CO-OH,

(6) -C \equiv N,

(V) $C_2\text{-}C_{10}$ alkenyl optionally substituted with one, two
25 or three substituents independently selected from the group
consisting of $C_1\text{-}C_3$ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N,
-CF $_3$, $C_1\text{-}C_6$ alkoxy, -O-phenyl, and -NR $_{1-a}$ R $_{1-b}$,

(VI) $C_2\text{-}C_{10}$ alkynyl optionally substituted with one,
two or three substituents independently selected from the group
30 consisting of $C_1\text{-}C_3$ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N,
-CF $_3$, $C_1\text{-}C_6$ alkoxy, -O-phenyl, and -NR $_{1-a}$ R $_{1-b}$,

(VII) -($C_1\text{-}C_6$ alkyl)-O-($C_1\text{-}C_6$ alkyl)-OH,

(VIII) -CH $_2$ -NH-CH $_2$ -CH(-O-CH $_2$ -CH $_3$) $_2$,

(IX) -(CH $_2$) $_{0-6}$ -C(=NR $_{1-a}$)(NR $_{1-a}$ R $_{1-b}$);

where R_N is

(I) $R_{N-1}-X_N-$ where X_N is $-\text{CO}-$, and where R_{N-1} is selected from the group consisting of:

(A) phenyl, which is optionally substituted with one,
5 two or three of the following substituents which can be the same or different and are:

(1) $\text{C}_1\text{-C}_6$ alkyl, optionally substituted with one,
two or three substituents selected from the group consisting of
 $\text{C}_1\text{-C}_3$ alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1\text{-C}_3$
10 alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

wherein R_{1-a} and R_{1-b} at each occurrence are independently H or $\text{C}_1\text{-C}_6$ alkyl,

(2) $-\text{OH}$,

(3) $-\text{NO}_2$,

15 (4) $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$,

(5) $-\text{CO}_2\text{H}$,

(6) $-\text{C}\equiv\text{N}$,

(7) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{N-2}\text{R}_{N-3}$ where R_{N-2} and R_{N-3} are the same or different and are selected from the group consisting of:

20 (a) $-\text{H}$,

(b) $-\text{C}_1\text{-C}_8$ alkyl optionally substituted with one substituent selected from the group consisting of:

(i) $-\text{OH}$,

(ii) $-\text{NH}_2$,

25 (iii) phenyl,

(c) $-\text{C}_1\text{-C}_8$ alkyl optionally substituted with 1, 2, or 3 groups that are independently $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, or $-\text{I}$,

(d) $-\text{C}_3\text{-C}_8$ cycloalkyl,

(e) $-(\text{C}_1\text{-C}_2 \text{ alkyl})-(\text{C}_3\text{-C}_8 \text{ cycloalkyl})$,

30 (f) $-(\text{C}_1\text{-C}_6 \text{ alkyl})-\text{O}-(\text{C}_1\text{-C}_3 \text{ alkyl})$,

(g) $-\text{C}_2\text{-C}_6$ alkenyl,

(h) $-\text{C}_2\text{-C}_6$ alkynyl,

(i) $-\text{C}_1\text{-C}_6$ alkyl chain with one double bond and one triple bond,

(j) $-R_{1-aryl}$, wherein R_{1-aryl} at each occurrence is independently phenyl, naphthyl, indanyl, indenyl, dihydronaphthyl, or tetralinyl each of which is optionally substituted with 1, 2, 3, or 4 groups that are independently:

5 (i) C_1-C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

10 (ii) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

15 (iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(iv) $-F$, Cl , $-Br$ and $-I$,

(v) $-C_1-C_6$ alkoxy optionally substituted with 1, 2, or 3 $-F$,

20 (vi) $-NR_{N-2}R_{N-3}$,

(vii) $-OH$,

(viii) $-C\equiv N$,

25 (ix) C_3-C_7 cycloalkyl, optionally substituted with 1, 2, or 3 groups that are selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(x) $-CO-(C_1-C_4 \text{ alkyl})$,

(xi) $-SO_2-NR_{1-a}R_{1-b}$,

(xii) $-CO-NR_{1-a}R_{1-b}$, or

30 (xiii) $-SO_2-(C_1-C_4 \text{ alkyl})$,

(k) $-R_{1-heteroaryl}$, wherein $R_{1-heteroaryl}$ at each occurrence is independently selected from the group consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl,

quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl,
 isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indolizinyl,
 indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl,
 furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl,
 5 triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl,
 isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-
 carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl,
 isoindolinyl, isobenzotetrahydrofuranlyl,
 isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl,
 10 pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl,
 purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl,
 pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
 dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
 dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl,
 15 coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-
 oxide, tetrahydroquinolinyl, dihydroquinolinyl,
 dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
 dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl,
 benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
 20 pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,
 indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide,
 quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-
 oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
 thiazolyl N-oxide, indolizinyl N-oxide, indazolyl N-oxide,
 25 benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
 oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
 oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and
 benzothiopyranyl S,S-dioxide,

where the R₁-heteroaryl group is optionally
 30 substituted with 1, 2, 3, or 4 groups that are independently:

(i) C₁-C₆ alkyl optionally
 substituted with 1, 2, or 3 groups independently selected from
 the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH,
 -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C₁-C₃ alkoxy,

(ii) C₂-C₆ alkenyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, or -NR_{1-a}R_{1-b},

(iii) C₂-C₆ alkynyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, or -NR_{1-a}R_{1-b},

(iv) -F, -Cl, -Br and -I,

(v) -C₁-C₆ alkoxy optionally substituted with one, two, or three -F,

(vi) -(CH₂)₀₋₄-NR_{N-2}R_{N-3},

(vii) -OH,

(viii) -C≡N,

(ix) (CH₂)₀₋₄-C₃-C₇ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(x) (CH₂)₀₋₄-CO-(C₁-C₆ alkyl),

(xi) (CH₂)₀₋₄-SO₂-NR_{N-2}R_{N-3},

(xii) (CH₂)₀₋₄-CO-NR_{N-2}R_{N-3},

(xiii) (CH₂)₀₋₄-SO₂-(C₁-C₆ alkyl),

(xiv) (CH₂)₀₋₄-N(R_{N-2})-SO₂-, and

(xv) (CH₂)₀₋₄-N(R_{N-2})-C(O)-,

(1) -R₁-heterocycle, wherein

R₁-heterocycle at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl, dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl, dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the R_1 -heterocycle group is bonded by any atom of the parent R_1 -heterocycle group substituted by hydrogen such that the new bond to the R_1 -heterocycle group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(a) C_1 - C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1 - C_3 alkyl, halogen, -OH, -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1 - C_3 alkoxy,

(b) C_2 - C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(c) C_2 - C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(d) halogen,

(e) C_1 - C_6 alkoxy,

(f) $-C_1$ - C_6 alkoxy optionally substituted with one, two, or three -F,

(g) $-NR_{N-2}R_{N-3}$,

(h) -OH,

(i) $-C\equiv N$,

(j) $(CH_2)_{0-4}$ -(C_3 - C_8 cycloalkyl), optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(k) $-(CH_2)_{0-4}-CO-(C_1-C_4$ alkyl),

(l) $-(CH_2)_{0-4}-SO_2-NR_{1-a}R_{1-b}$,

(m) $-(CH_2)_{0-4}-CO-NR_{1-a}R_{1-b}$,

(n) $-(CH_2)_{0-4}-SO_2-(C_1-C_6$ alkyl), and

(o) =O,

(p) $-(CH_2)_{0-4}-N(R_{N-2})-SO_2-$

(q) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{C}(\text{O})-$

(8) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_{12} \text{ alkyl}),$

(9) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkenyl}),$

(10) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkynyl}),$

5

(11) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl}),$

(12) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-aryl},$

(13) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl},$

(14) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle},$

(15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$ wherein $\text{R}_{\text{N}-4}$ is selected from the group consisting of phenyl, morpholinyl, thiomorpholinyl, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl, thienyl, pyrazolyl, pyridyl N-oxide, oxazolyl, thiazolyl, imidazolyl, and pyrrolidinyl where each group is optionally substituted with one, two, three, or four groups that are independently C_1-C_6 alkyl,

(16) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$ where $\text{R}_{\text{N}-5}$ is selected from the group consisting of:

(a) C_1-C_6 alkyl,

20

(b) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-aryl}),$

(c) C_2-C_6 alkenyl,

(d) C_2-C_6 alkynyl,

(e) $-(\text{CH}_2)_{0-2}-\text{C}_3-\text{C}_8 \text{ cycloalkyl},$

(f) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heteroaryl}),$ and

25

(g) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heterocycle}),$

(17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3},$

(18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl}),$

(19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl}),$

(20) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3-\text{C}_8 \text{ cycloalkyl}),$

30

(21) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{O}-\text{R}_{\text{N}-5},$

(22) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2,$

(23) $-(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2,$

(24) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2},$

(25) $-(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3},$

35

(26) $-(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4},$

(27) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl}),$

(28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$ wherein

R_{100} at each occurrence is independently $-\text{H}$ or C_1-C_4 alkyl,

5 (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2,$

(30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2,$

(31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5}),$

(32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH},$

(33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5}),$

10 (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl optionally substituted with one, two, three, four, or five of } -\text{F}),$

(35) C_3-C_8 cycloalkyl,

(36) C_2-C_6 alkenyl optionally substituted with C_1-C_3 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy,
15 or $-\text{NR}_{1-a}\text{R}_{1-b},$

(37) C_2-C_6 alkynyl optionally substituted with C_1-C_3 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy,
or $-\text{NR}_{1-a}\text{R}_{1-b},$

(38) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2},$

20 (39) $-(\text{CH}_2)_{1-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl}),$

(B) $-\text{R}_{\text{N-heteroaryl}}$ where $\text{R}_{\text{N-heteroaryl}}$ is selected from the group consisting of pyridinyl, indolyl, indolinyl, isoindolyl, imidazolyl, isoxazolyl, oxazolyl, thiazolyl, indolizinyll and isochromanyl,

25 where the $\text{R}_{\text{N-heteroaryl}}$ group is bonded by any atom of the parent $\text{R}_{\text{N-heteroaryl}}$ group substituted by hydrogen such that the new bond to the $\text{R}_{\text{N-heteroaryl}}$ group replaces the hydrogen atom and its bond, where heteroaryl is optionally substituted with one, two, three, or four of:

30 (1) C_1-C_6 alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C_1-C_3 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b},$

(2) $-\text{OH},$

- (3) $-\text{NO}_2$,
- (4) $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$,
- (5) $-\text{CO}_2\text{H}$,
- (6) $-\text{C}\equiv\text{N}$,
- 5 (7) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
- (8) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
- (9) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkenyl})$,
- (10) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkynyl})$,
- (11) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
- 10 (12) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-aryl}$,
- (13) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl}$,
- (14) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle}$,
- (15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$,
- (16) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$,
- 15 (17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
- (18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl})$,
- (19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
- (20) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
- (21) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$,
- 20 (22) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (23) $-(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (24) $-(\text{CH}_2)_{0-4}-\text{N}(-\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2}$,
- (25) $-(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
- (26) $-(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4}$,
- 25 (27) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
- (28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$,
- (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
- 30 (32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
- (33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
- (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl optionally substituted with one, two, three, four, or five of } -\text{F})$,
- (35) $\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,

(36) C₂-C₆ alkenyl optionally substituted with C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, or -NR_{1-a}R_{1-b},

(37) C₂-C₆ alkynyl optionally substituted with C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, or -NR_{1-a}R_{1-b},

(38) -(CH₂)₀₋₄-N(-H or R_{N-5})-SO₂-R_{N-2},

(39) -(CH₂)₁₋₄-C₃-C₈ cycloalkyl,

(C) R_N-aryl-W-R_N-aryl,

(D) R_N-aryl-W-R_N-heteroaryl,

(E) R_N-aryl-W-R₁-heterocycle,

(F) R_N-heteroaryl-W-R_N-aryl,

(G) R_N-heteroaryl-W-R_N-heteroaryl,

(H) R_N-heteroaryl-W-R_{N-1}-heterocycle,

(I) R_N-heterocycle-W-R_N-aryl,

(J) R_N-heterocycle-W-R_N-heteroaryl,

(K) R_N-heterocycle-W-R_{N-1}-heterocycle,

where W is

(7) -(CH₂)₁₋₄-,

(8) -O-,

(9) -S(O)₀₋₂-,

(10) -N(R_{N-5})-,

(11) -CO-; or

(12) a bond;

(II) -CO-(C₁-C₆ alkyl)-M-(C₁-C₆ alkyl), where M is S, SO or SO₂, and wherein each alkyl is unsubstituted or substituted with one, two, or three of substituents independently selected from the group consisting of:

(A) -NH-CO-(C₁-C₆ alkyl),

(B) -NH-CO-O-R_{N-8},

(C) -NR_{N-2}R_{N-3};

where R₁ is

-(CH₂)_{n₁}-phenyl, where n₁ is zero or one, and which is optionally substituted with one, two, three or four of the following substituents on the phenyl ring:

(A) C₁-C₆ alkyl optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

5 (B) C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

10 (C) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(D) -F, Cl, -Br or -I,

15 (F) -C₁-C₆ alkoxy optionally substituted with one, two or three of -F,

(G) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(H) -OH,

(I) -C≡N,

20 (J) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(K) -CO-(C₁-C₄ alkyl),

25 (L) -SO₂-NR_{1-a}R_{1-b},

(M) -CO-NR_{1-a}R_{1-b},

(N) -SO₂-(C₁-C₄ alkyl); and

where R₂ is

30 (I) -(Z)-C₁-C₆ alkyl, where Z is a bond, -C(O), -CO₂- or -SO₂-, wherein the alkyl group is optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, C₁-C₇ alkyl (optionally substituted with C₁-C₃ alkyl and C₁-C₃ alkoxy), -F, -Cl, -Br, -I, -OH, -SH,

-C≡N, -CF₃, C₁-C₃ alkoxy, -NR_{1-a}R_{1-b} where R_{1-a} and R_{1-b} are independently -H or C₁-C₆ alkyl, and -OC=O NR_{1-a}R_{1-b},

(II) -(Z)-CH₂-S(O)₀₋₂-(C₁-C₆ alkyl),

(III) -(Z)-CH₂-CH₂-S(O)₀₋₂-(C₁-C₆ alkyl),

5 (IV) -(Z)-C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(V) -(Z)-C₂-C₆ alkynyl with one or two triple bonds,
10 optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(VI) -(Z)-(CH₂)_{n1}-(R_{1-aryl}), where Z is a bond, CO, CO₂ or SO₂, where n₁ is zero or one and where R_{1-aryl} is phenyl, 1-naphthyl, 2-naphthyl and indanyl, indenyl, dihydronaphthalyl, or
15 tetralinyl optionally substituted with one, two, three or four of the following substituents on the aryl ring:

(A) C₁-C₆ alkyl optionally substituted with one, two or three substituents selected from the group consisting of
20 C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(B) C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -
25 CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(C) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -
CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

30 (D) -F, Cl, -Br or -I,

(F) -C₁-C₆ alkoxy optionally substituted with one, two or three of -F,

(G) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(H) -OH,

(I) -C≡N,

(J) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(K) -CO-(C₁-C₄ alkyl),(L) -SO₂-NR_{1-a}R_{1-b},(M) -CO-NR_{1-a}R_{1-b},

10 (N) -SO₂-(C₁-C₄ alkyl),

(VII) -(Z)-(CH₂)_{n1}-(R_{1-heteroaryl}) where n₁ is as defined above and where R_{1-heteroaryl} is selected from the group consisting of:

pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl,
 15 indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl,
 20 triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl,
 25 pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl, dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl, dihydrobenzisothiazinyl, benzopyranlyl, benzothiopyranlyl,
 30 coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-oxide, tetrahydroquinolinyl, dihydroquinolinyl, dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl, dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl, benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
 35 pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,

indolyl N-oxide, indoliny N-oxide, isoquinolyl N-oxide,
quinazoliny N-oxide, quinoxaliny N-oxide, phthalaziny N-
oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
thiazolyl N-oxide, indoliziny N-oxide, indazolyl N-oxide,
5 benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide,
benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is bonded to $-(CH_2)_{n1}-$ by any ring atom
10 of the parent R_N -heteroaryl group substituted by hydrogen such that
the new bond to the R_1 -heteroaryl group replaces the hydrogen atom
and its bond, where heteroaryl is optionally substituted with
one, two, three or four of:

(1) C_1-C_6 alkyl optionally substituted with one,
15 two or three substituents selected from the group consisting of
 C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$,
 $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(2) C_2-C_6 alkenyl with one or two double bonds,
optionally substituted with one, two or three substituents
20 selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-$
 CF_3 , C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(3) C_2-C_6 alkynyl with one or two triple bonds,
optionally substituted with one, two or three substituents
selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-$
25 CF_3 , C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(4) $-F$, Cl , $-Br$ or $-I$,

(6) $-C_1-C_6$ alkoxy optionally substituted with
one, two, or three of $-F$,

(7) $-NR_{N-2}R_{N-3}$ where R_{N-2} and R_{N-3} are as defined
30 below,

(8) $-OH$,

(9) $-C\equiv N$,

(10) C_3-C_7 cycloalkyl, optionally substituted
with one, two or three substituents selected from the group

consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(11) -CO-(C₁-C₄ alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

5 (13) -CO-NR_{1-a}R_{1-b}, or

(14) -SO₂-(C₁-C₄ alkyl), with the proviso that when n₁ is zero R_{1-heteroaryl} is not bonded to the carbon chain by nitrogen, or

(VIII) -(Z)-(CH₂)_{n1}-(R_{1-heterocycle}) where n₁ is as defined
 10 above and R_{1-heterocycle} is selected from the group consisting of:
 morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide,
 thiomorpholinyl S,S-dioxide, piperazinyl,
 homopiperazinyl, pyrrolidinyl, pyrrolinyl,
 tetrahydropyranyl, piperidinyl, tetrahydrofuranyl,
 15 tetrahydrothienyl, homopiperidinyl, homomorpholinyl,
 homomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide,
 oxazolidinonyl, dihydropyrazolyl,
 dihydropyrrolyl dihydropyrazinyl dihydropyridinyl
 dihydropyrimidinyl, dihydrofuryl, dihydropyranyl,
 20 tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, homothiomorpholinyl S-oxide,

where the R_{1-heterocycle} group is bonded by any atom of the parent R_{1-heterocycle} group substituted by hydrogen such that the new bond to the R_{1-heterocycle} group replaces the hydrogen atom and its
 25 bond, where heterocycle is optionally substituted with one, two, three or four:

(1) C₁-C₆ alkyl optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -
 30 CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(2) C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(3) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

5 (4) -F, Cl, -Br, or -I,

(5) C₁-C₆ alkoxy,

(6) -C₁-C₆ alkoxy optionally substituted with one, two, or three -F,

10 (7) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(8) -OH,

(9) -C≡N,

15 (10) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(11) -CO-(C₁-C₄ alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

(13) -CO-NR_{1-a}R_{1-b},

20 (14) -SO₂-(C₁-C₄ alkyl),

(15) =O, with the proviso that when n₁ is zero R_{1-heterocycle} is not bonded to the carbon chain by nitrogen; and

where R₂₀ is H or C₁₋₆ alkyl or alkenyl.

25 In another preferred embodiment relative to formula II, R_c is -(CR_{C-x}R_{C-y})₀₋₄-R_{C-aryl} where R_{C-x} and R_{C-y} are independently selected from the group consisting of

-H,

C₁-C₄ alkyl optionally substituted with 1 or 2 -OH,

30 C₁-C₄ alkoxy optionally substituted with 1, 2, or 3 halogen,

-(CH₂)₀₋₄-C₃-C₈ cycloalkyl,

C₂-C₆ alkenyl containing one or two double bonds,

C₂-C₆ alkynyl containing one or two triple bonds, and

phenyl,

or

R_{C-x} and R_{C-y} are taken together with the carbon to which they are attached to form a carbocycle of three, four, five, six or seven carbon atoms, where one carbon atom is optionally replaced by a group selected from -O-, -S-, -SO₂-, -NR_{N-2}- and R_{C-aryl} , wherein

R_{C-aryl} is phenyl, which is optionally substituted with 1, 2, or 3 groups that are independently:

(1) C₁-C₆ alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(2) -OH,

(3) -NO₂,

(4) halogen,

(5) -CO₂H,

(6) -C≡N.

In yet another preferred embodiment, R_c is -(CR_{C-x}R_{C-y})₀₋₄-R_{C-aryl} where R_{C-x} and R_{C-y} are independently selected from the group consisting of

-H,

C₁-C₄ alkyl optionally substituted with 1 or 2 -OH,

C₁-C₄ alkoxy optionally substituted with 1, 2, or 3 halogen,

-(CH₂)₀₋₄-C₃-C₈ cycloalkyl,

C₂-C₆ alkenyl containing one or two double bonds,

C₂-C₆ alkynyl containing one or two triple bonds, and phenyl,

or

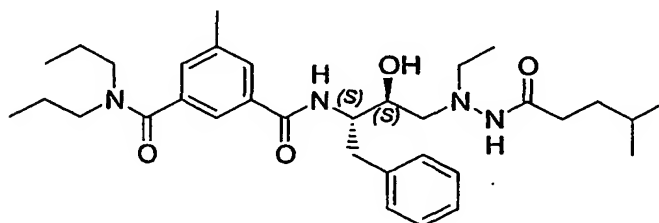
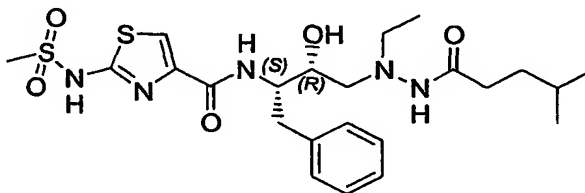
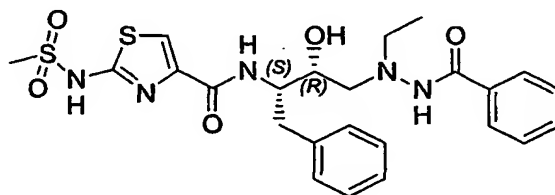
R_{C-x} and R_{C-y} are taken together with the carbon to which they are attached to form a carbocycle of three, four, five, six or seven carbon atoms, where one carbon atom is optionally replaced by a group selected from -O-, -S-, -SO₂-, -NR_{N-2}- and R_{C-aryl} , wherein

-(CR_{C-x}R_{C-y})₀₋₄-R_C-heteroaryl is selected from the group consisting of pyridinyl, indolyl, indolinyl, isoindolyl, imidazolyl, isoxazolyl, oxazolyl, thiazolyl, indolizinyl and isochromanlyl.

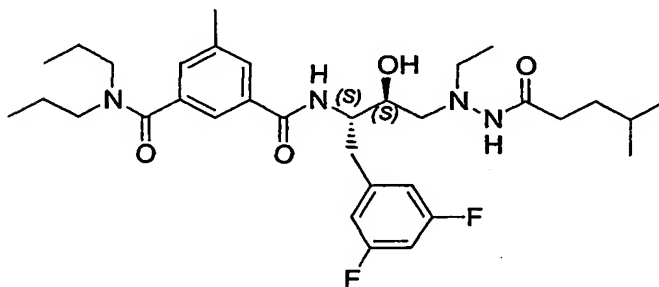
5 In another preferred embodiment relative to formula II, R_C is the optionally substituted -C₁-C₁₀ alkyl groups as described above.

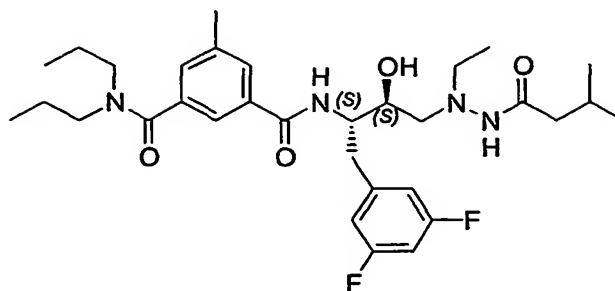
Preferred compounds of the formula II include, amongst others:

10

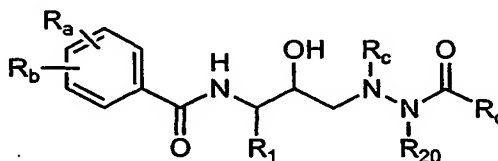


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In another preferred embodiment, the invention provides
5 compounds of formula III:



(III)

or a pharmaceutically acceptable salt thereof wherein

- 10 R_1 represents phenyl (C_1 - C_6)alkyl where the phenyl is optionally substituted with up to three groups independently selected from halogen, hydroxy, C_1 - C_2 alkyl, C_1 - C_2 alkoxy, amino, nitro, trifluoromethyl, cyano, mono(C_1 - C_2)alkylamino and di(C_1 - C_2)alkylamino;
- 15 R_a and R_b independently represent hydrogen, hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, C_3 - C_7 cycloalkyl, C_3 - C_7 cycloalkyl(C_1 - C_6)alkyl, C_3 - C_7 cycloalkyl(C_1 - C_6)alkoxy, halogen, cyano, amino, mono(C_1 - C_6)alkylamino, di(C_1 - C_6)alkylamino, mono- or di(C_1 - C_6)alkylaminosulfonyl, C_1 - C_6 alkyl sulfonylamino, C_2 - C_6
- 20 alkenyl, C_2 - C_6 alkynyl, trifluoromethyl, mono(C_1 - C_6)alkylaminocarbonyl, or di(C_1 - C_6)alkylaminocarbonyl and

provided that not both R_a and R_b are hydrogen simultaneously;

R_c represents hydrogen, or C_1 - C_6 alkyl, C_2 - C_6 alkenyl, or C_2 - C_6 alkynyl each of which is optionally substituted with
5 halogen, hydroxy, amino, cyano, or trifluoromethyl;

R_d represents

phenyl optionally substituted with hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, C_3 - C_7 cycloalkyl, C_3 - C_7 cycloalkyl(C_1 - C_6)alkyl, C_3 - C_7 cycloalkyl(C_1 - C_6)alkoxy, halogen,
10 cyano, amino, mono(C_1 - C_6)alkylamino, di(C_1 - C_6)alkylamino, mono- or di(C_1 - C_6)alkylaminosulfonyl, C_1 - C_6 alkyl sulfonylamino, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl;
or

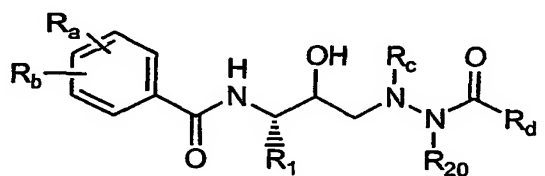
C_1 - C_6 alkyl optionally substituted with hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halogen, cyano, amino, mono(C_1 - C_6)alkylamino, di(C_1 - C_6)alkylamino, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, or trifluoromethyl; and
15

R_{20} represents hydrogen, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, or trifluoromethyl.

20 In accordance with this preferred embodiment, R_1 is benzyl where the phenyl is optionally substituted. Also preferably, the phenyl is substituted with one or two groups independently selected from halogen, hydroxy, C_1 - C_3 alkyl, amino, and trifluoromethyl. In another preferred embodiment, phenyl is
25 substituted with two groups independently selected from halogen, hydroxy, and trifluoromethyl. In an alternative preferred

embodiment, phenyl is disubstituted with halogen. Also preferably, R_1 is 3,5-difluorobenzyl. Preferably, R_a and R_b are different and R_b represents mono- or di(C_1 - C_6)alkylaminocarbonyl. Also preferably, R_d is phenyl optionally substituted with C_1 - C_3 alkyl, C_1 - C_3 alkoxy, amino, hydroxy, or halogen. In a further preferred embodiment, R_c is hydrogen or C_1 - C_4 alkyl. Also preferably, R_c is C_1 - C_3 alkyl. In yet another preferred embodiment, R_d is C_1 - C_6 lower alkyl and R_{20} is hydrogen.

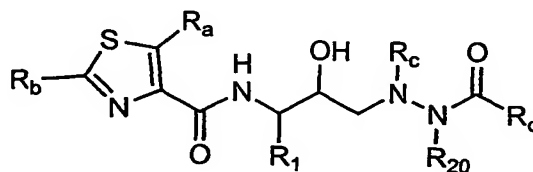
In another preferred embodiment, the invention provides for compounds of formula IV:



(IV)

where R_a , R_b , R_1 , R_c , R_{20} and R_d are defined above for this preferred embodiment. In another preferred embodiment, R_1 is benzyl where the phenyl is disubstituted with chloro or fluoro; R_c is C_1 - C_3 alkyl; R_d is C_1 - C_6 lower alkyl; R_{20} is hydrogen or C_1 - C_6 alkyl; and R_b is di(C_1 - C_6)alkylaminocarbonyl attached to the 3-position of the phenyl group.

In another preferred embodiment, the invention provides compounds of the formula V:



(V)

or a pharmaceutically acceptable salt thereof wherein

R₁ represents phenyl (C₁-C₆)alkyl where the phenyl is optionally substituted with up to three groups independently selected from halogen, hydroxy, C₁-C₂ alkyl, C₁-C₂ alkoxy, amino, nitro, trifluoromethyl, cyano, mono(C₁-C₂)alkylamino and di(C₁-C₂)alkylamino;

R_a and R_b independently represent hydrogen, hydroxy, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₃-C₇ cycloalkyl, C₃-C₇ cycloalkyl(C₁-C₆)alkyl, C₃-C₇ cycloalkyl(C₁-C₆)alkoxy, halogen, cyano, amino, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, mono- or di(C₁-C₆)alkylaminosulfonyl, C₁-C₆ alkyl sulfonylamino, C₂-C₆ alkenyl, C₂-C₆ alkynyl, trifluoromethyl, mono(C₁-C₆)alkylaminocarbonyl, or di(C₁-C₆)alkylaminocarbonyl and provided that not both R_a and R_b are hydrogen simultaneously;

R_c represents hydrogen, or C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl each of which is optionally substituted with halogen, hydroxy, amino, cyano, or trifluoromethyl;

R_d represents

phenyl optionally substituted with hydroxy, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₃-C₇ cycloalkyl, C₃-C₇ cycloalkyl(C₁-C₆)alkyl, C₃-C₇ cycloalkyl(C₁-C₆)alkoxy, halogen, cyano, amino, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, mono- or di(C₁-C₆)alkylaminosulfonyl,

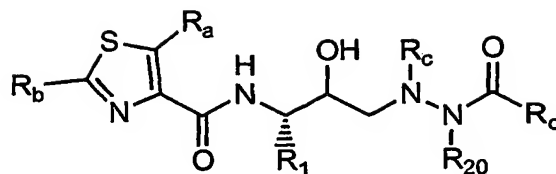
C₁-C₆ alkyl sulfonylamino, C₂-C₆ alkenyl, C₂-C₆ alkynyl;

or

C₁-C₆ alkyl optionally substituted with hydroxy, C₁-C₆ alkyl, C₁-C₆ alkoxy, halogen, cyano, amino, mono(C₁-
5 C₆)alkylamino, di(C₁-C₆)alkylamino, C₂-C₆ alkenyl, C₂-C₆ alkynyl, or trifluoromethyl; and

R₂₀ represents hydrogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, or trifluoromethyl.

In accordance with this preferred embodiment, R₁ is benzyl where
10 the phenyl is optionally substituted. Preferably, the phenyl is substituted with one or two groups independently selected from halogen, hydroxy, C₁-C₃ alkyl, amino, and trifluoromethyl. Also preferably, phenyl is substituted with two groups independently selected from halogen, hydroxy, and trifluoromethyl. Also
15 preferably, phenyl is disubstituted with halogen. In another preferred embodiment, R₁ is 3,5-difluorobenzyl. Also preferably, R_a and R_b are different and R_b represents C₁-C₆)alkylsulfonylamino. Preferably, R_d is phenyl optionally substituted with C₁-C₃ alkyl, C₁-C₃ alkoxy, amino, hydroxy, or
20 halogen. Preferably, R_c is hydrogen or C₁-C₄ alkyl. Preferably, R_c is C₁-C₃ alkyl. Also preferably, R_d is C₁-C₆ lower alkyl and R₂₀ is hydrogen. Further in accordance with this preferred embodiment, the invention provides compounds of the formula VI:



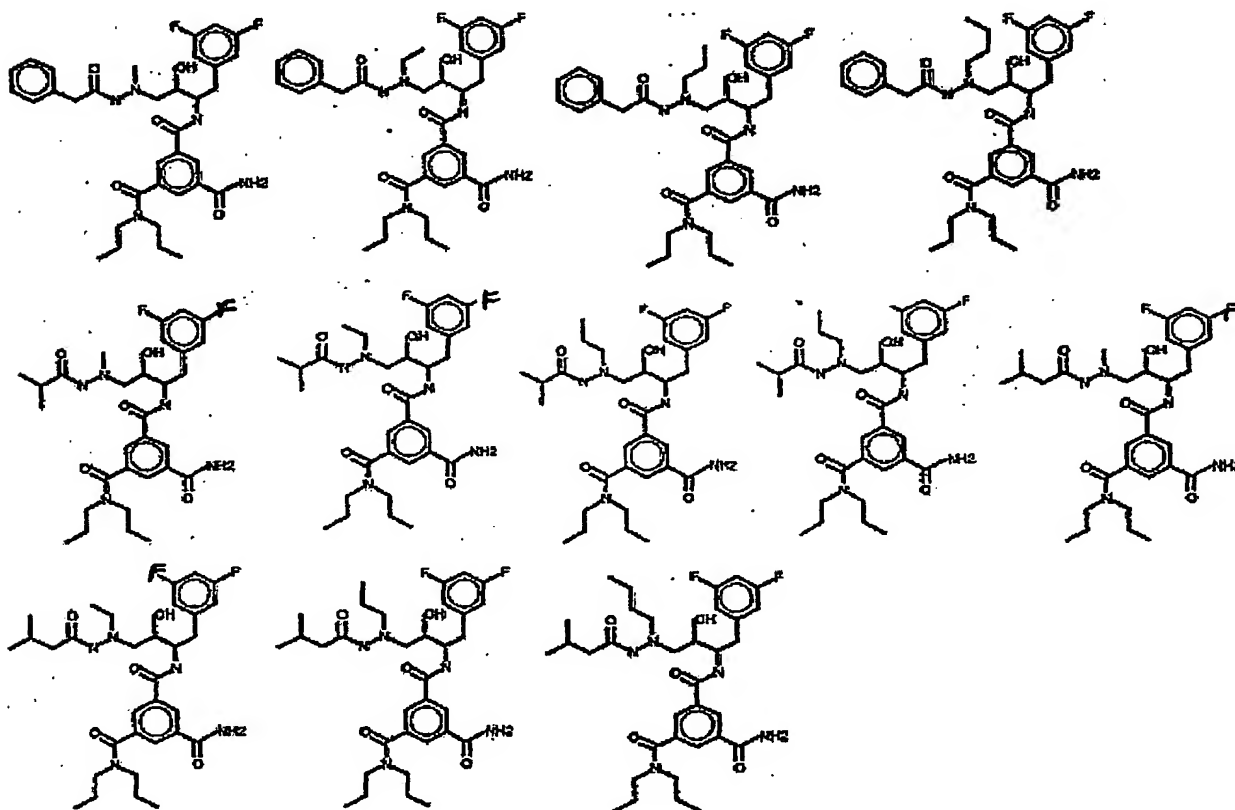
(VI)

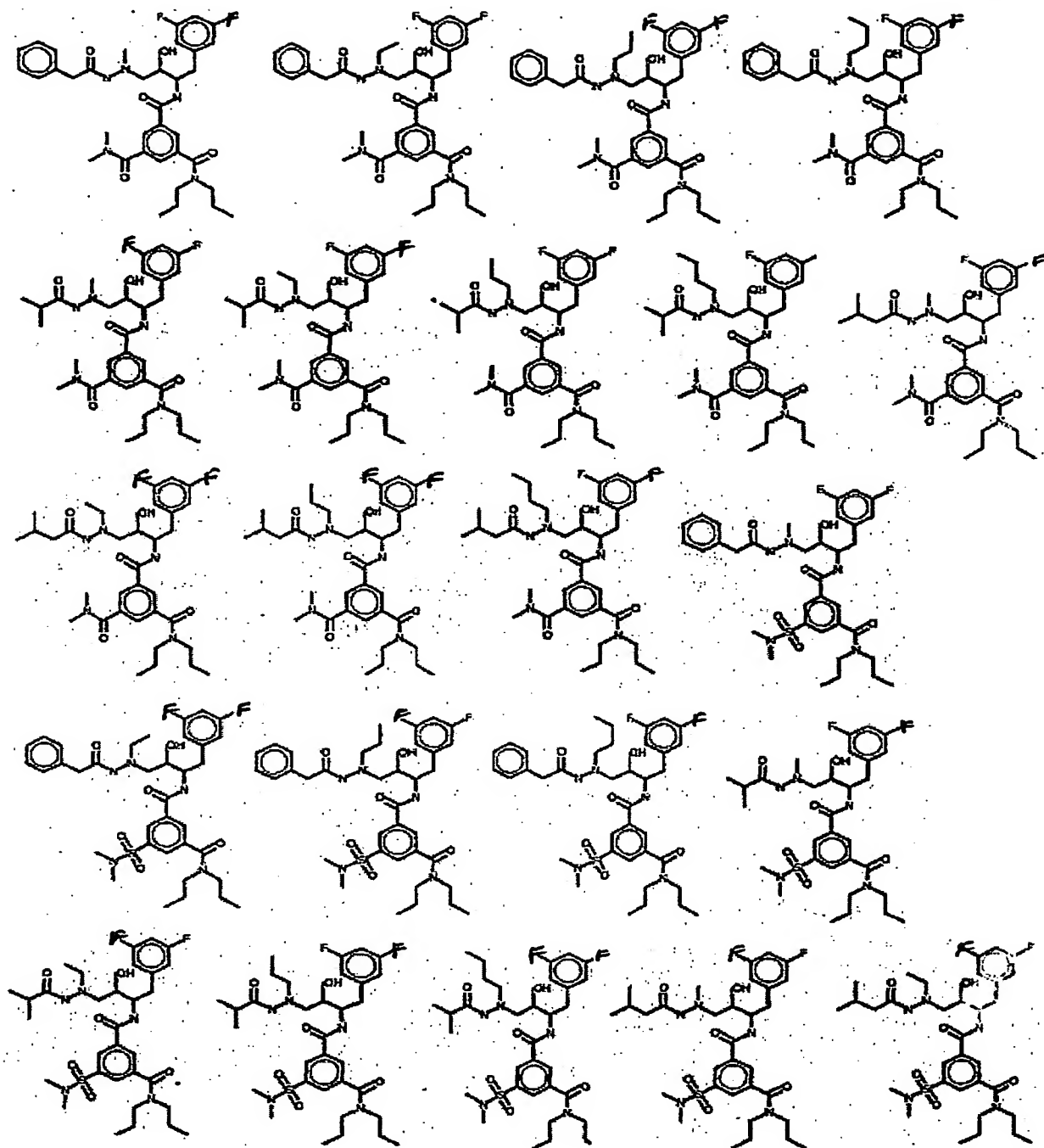
where R_a , R_b , R_1 , R_c , R_{20} and R_d are defined above for this

preferred embodiment. In another preferred embodiment, R_1 is

5 benzyl where the phenyl is disubstituted with chloro or fluoro; R_c is C_1 - C_3 alkyl; R_d is C_1 - C_6 lower alkyl; R_{20} is hydrogen or C_1 - C_6 alkyl; and R_b is alkylsulfonylamino attached to the 2-position of the thiazolyl group.

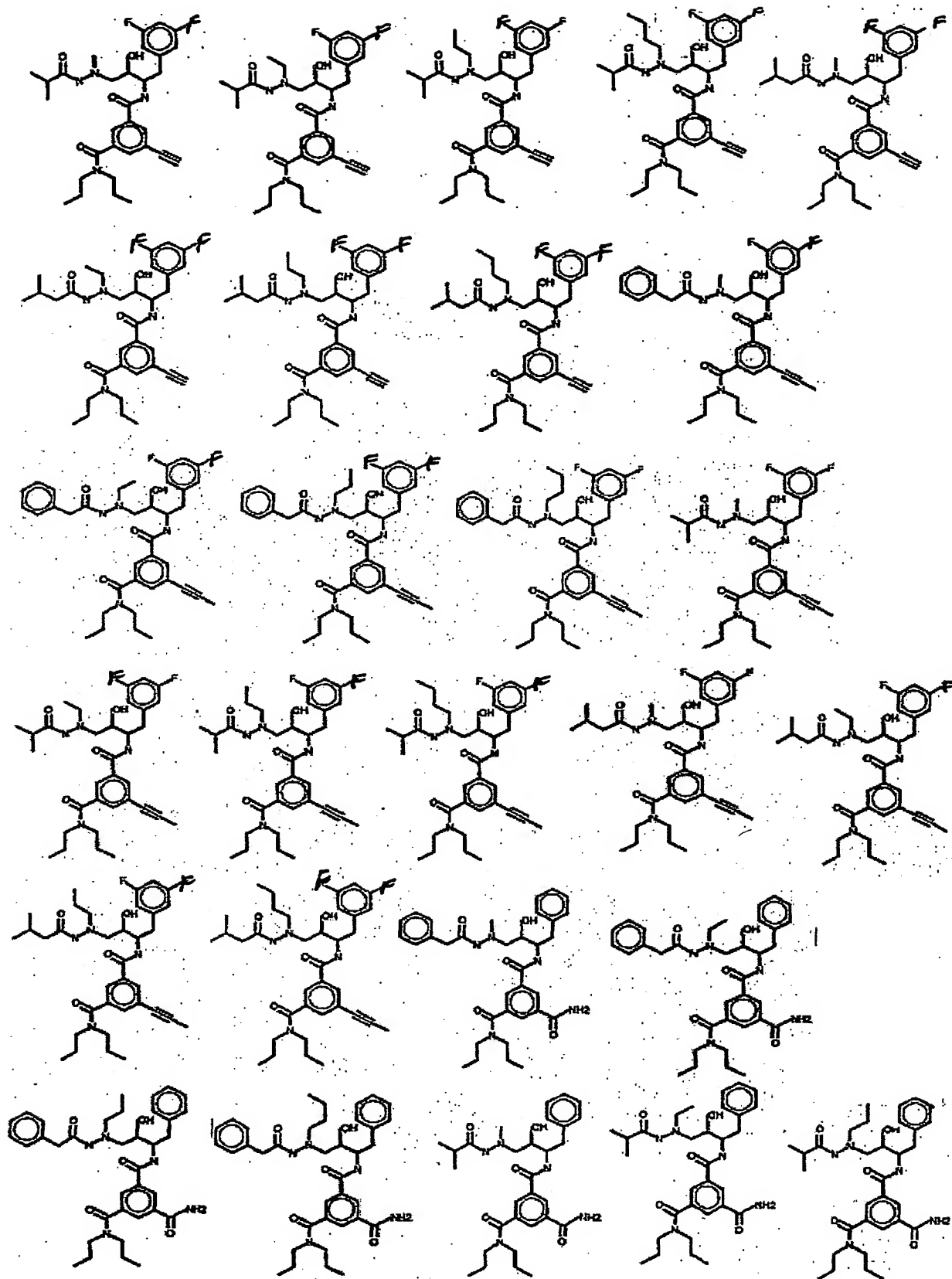
Preferred compounds of the invention include:

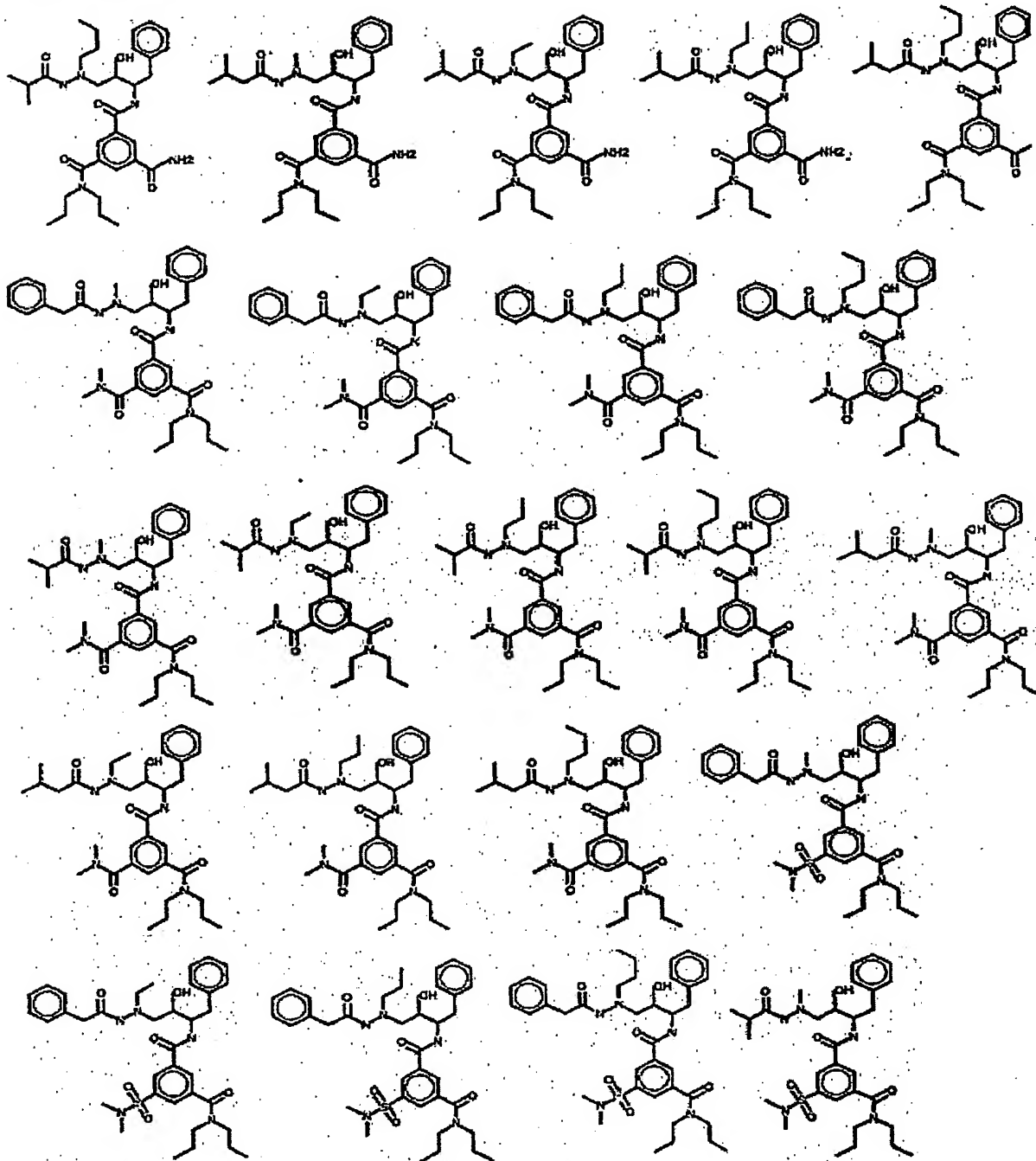


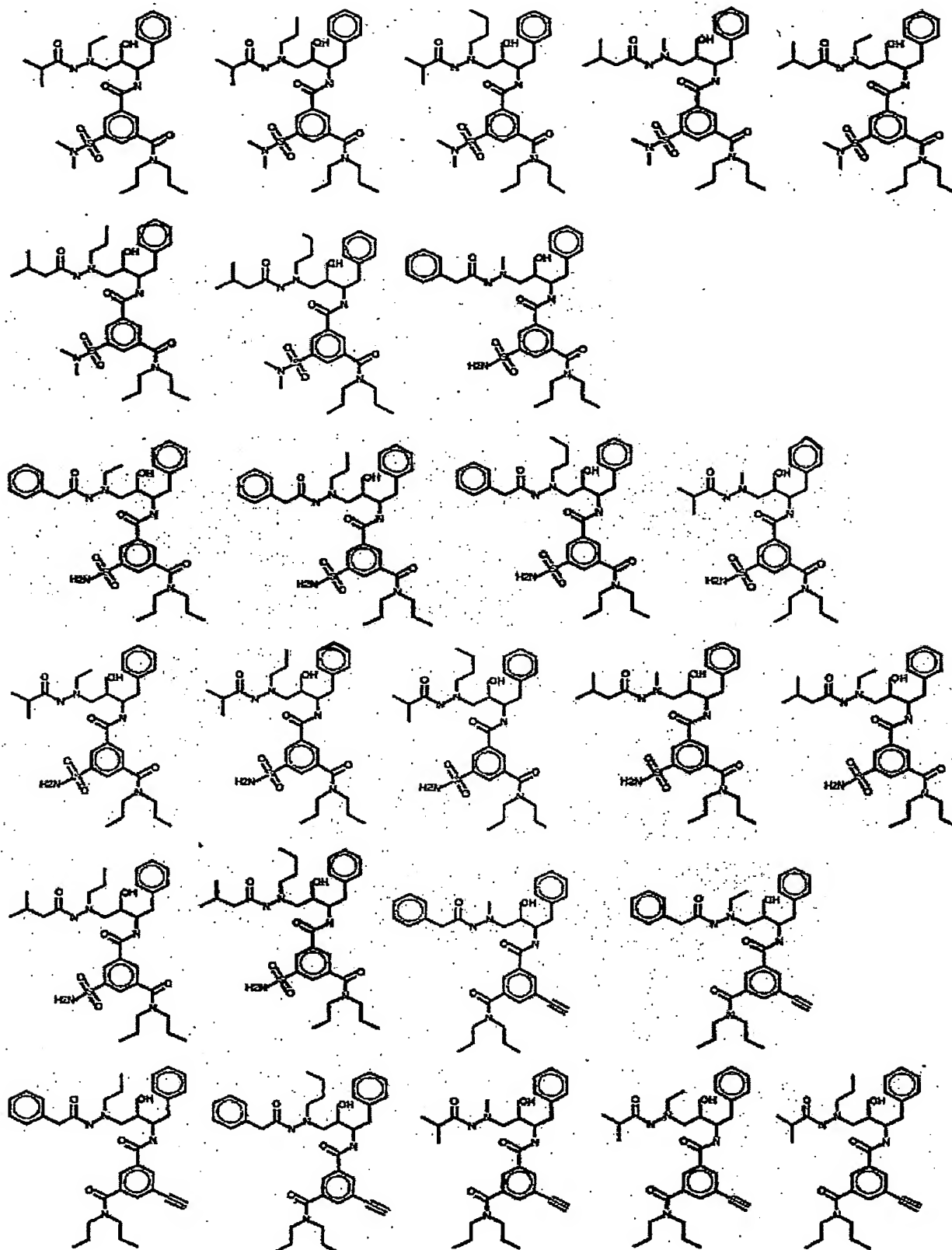


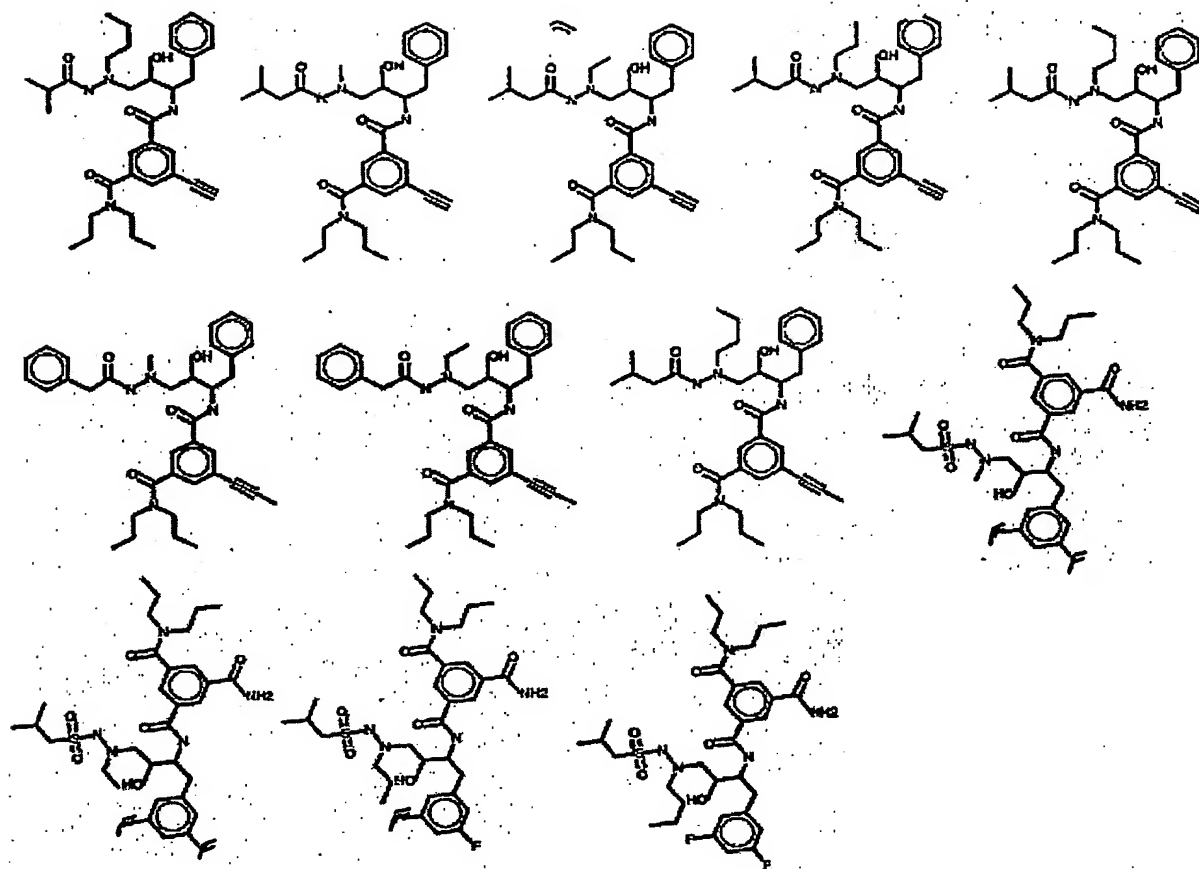
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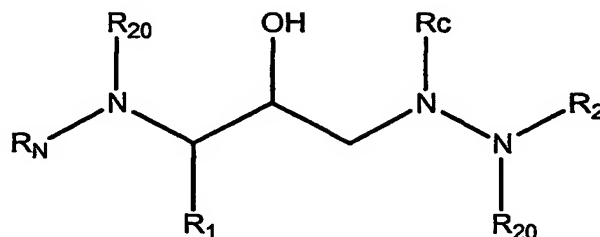


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In another embodiment, the invention provides a method of treating a patient who has, or in preventing a patient from getting, a disease or condition selected from the group consisting of Alzheimer's disease, for helping prevent or delay the onset of Alzheimer's disease, for treating patients with mild cognitive impairment (MCI) and preventing or delaying the onset of Alzheimer's disease in those who would progress from MCI to AD, for treating Down's syndrome, for treating humans who have Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, for treating cerebral amyloid angiopathy and preventing its potential consequences, i.e. single and recurrent lobar hemorrhages, for treating other degenerative dementias, including dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's disease, dementia associated with progressive supranuclear palsy, dementia associated with cortical basal degeneration, diffuse Lewy body

type of Alzheimer's disease and who is in need of such treatment which comprises administration of a therapeutically effective amount of a compound selected from the group consisting of an aza hydroxylated ethyl amine of the formula II:

5



(II)

or a pharmaceutically acceptable salt thereof,

10 where Rc is

(I) -C₁-C₁₀ alkyl optionally substituted with one, two or three groups independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, -NR_{1-a}R_{1-b}, -OC(=O)NR_{1-a}R_{1-b}, -S(=O)₀₋₂R_{1-a}, -NR_{1-a}C(=O)NR_{1-a}R_{1-b},
15 -C(=O)NR_{1-a}R_{1-b}, and -S(=O)₂NR_{1-a}R_{1-b} wherein

R_{1-a} and R_{1-b} at each occurrence are independently H or C₁-C₆ alkyl,

(II) -(CH₂)₀₋₃-(C₃-C₈) cycloalkyl where cycloalkyl can be optionally substituted with one, two or three substituents
20 independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, -CO₂H, -CO₂-(C₁-C₄ alkyl), and -NR_{1-a}R_{1-b}

(III) -(CR_{C-x}R_{C-y})₀₋₄-R_{C-ary1} where R_{C-x} and R_{C-y} are independently selected from the group consisting of

25 -H,
C₁-C₄ alkyl optionally substituted with 1 or 2 -OH,
C₁-C₄ alkoxy optionally substituted with 1, 2, or 3 halogen,

- (CH₂)₀₋₄-C₃-C₈ cycloalkyl,
30 C₂-C₆ alkenyl containing one or two double bonds,
C₂-C₆ alkynyl containing one or two triple bonds, and

phenyl,

or

R_{C-x} and R_{C-y} are taken together with the carbon to which they are attached to form a carbocycle of three, four, five, six or seven carbon atoms, where one carbon atom is optionally replaced by a group selected from -O-, -S-, -SO₂-, -NR_{N-2}- and R_{C-aryl} , wherein

R_{C-aryl} is phenyl, which is optionally substituted with 1, 2, or 3 groups that are independently:

(1) C₁-C₆ alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(2) -OH,

(3) -NO₂,

(4) halogen,

(5) -CO₂H,

(6) -C≡N,

(7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3} where

R_{N-2} and R_{N-3} are independently selected from the group consisting of:

(a) -H,

(b) -C₁-C₆ alkyl optionally substituted with one substituent selected from the group consisting of:

(i) -OH, and

(ii) -NH₂,

(c) -C₁-C₆ alkyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -Br, -I, or OH,

(d) -C₃-C₇ cycloalkyl,

(e) -(C₁-C₂ alkyl)-(C₃-C₇ cycloalkyl),

(f) -(C₁-C₆ alkyl)-O-(C₁-C₃ alkyl),

(g) -C₂-C₆ alkenyl

(h) -C₂-C₆ alkynyl

(i) $-C_1-C_6$ alkyl chain with one double bond and one triple bond,

(j) $-R_{1-aryl}$ wherein R_{1-aryl} at each occurrence is independently phenyl, naphthyl, indanyl, indenyl, dihydronaphthyl, or tetralinyl each of which is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(i) C_1-C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

(ii) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(iv) $-F$, Cl , $-Br$ and $-I$,

(v) $-C_1-C_6$ alkoxy optionally substituted with 1, 2, or 3 $-F$,

(vi) $-NR_{N-2}R_{N-3}$,

(vii) $-OH$,

(viii) $-C\equiv N$,

(ix) C_3-C_7 cycloalkyl, optionally substituted with 1, 2, or 3 groups that are selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(x) $-CO-(C_1-C_4 \text{ alkyl})$,

(xi) $-SO_2-NR_{1-a}R_{1-b}$,

(xii) $-CO-NR_{1-a}R_{1-b}$, or

(xiii) $-SO_2-(C_1-C_4 \text{ alkyl})$,

(k) $-R_{1-heteroaryl}$ wherein $R_{1-heteroaryl}$ at each occurrence is independently selected from the group

consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl, dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl, dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl, coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-oxide, tetrahydroquinolinyl, dihydroquinolinyl, dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl, dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl, benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide, pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide, indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide, quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide, thiazolyl N-oxide, indoliziny, indazolyl N-oxide, benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and benzothiopyranyl S,S-dioxide,

where the R₁-heteroaryl group is optionally substituted with 1, 2, 3, or 4 groups that are independently:

- (i) C_1-C_6 alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,
- 5 (ii) C_2-C_6 alkenyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,
- (iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,
- 10 (iv) -F, -Cl, -Br and -I,
- (v) $-C_1-C_6$ alkoxy optionally substituted with one, two, or three -F,
- 15 (vi) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,
- (vii) -OH,
- (viii) $-C\equiv N$,
- (ix) $(CH_2)_{0-4}-C_3-C_7$ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,
- 20 (x) $(CH_2)_{0-4}-CO-(C_1-C_6 \text{ alkyl})$,
- (xi) $(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,
- (xii) $(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,
- 25 (xiii) $(CH_2)_{0-4}-SO_2-(C_1-C_6 \text{ alkyl})$,
- (xiv) $(CH_2)_{0-4}-N(R_{N-2})-SO_2-$, and
- (xv) $(CH_2)_{0-4}-N(R_{N-2})-C(O)-$,
- 30 (8) $-(CH_2)_{0-4}-CO-(C_1-C_{12} \text{ alkyl})$,
- (9) $-(CH_2)_{0-4}-CO-(C_2-C_{12} \text{ alkenyl})$,
- (10) $-(CH_2)_{0-4}-CO-(C_2-C_{12} \text{ alkynyl})$,
- (11) $-(CH_2)_{0-4}-CO-(CH_2)_{0-4}-(C_3-C_7 \text{ cycloalkyl})$,
- (12) $-(CH_2)_{0-4}-CO-R_{1-aryl}$,

(13) - (CH₂)₀₋₄-CO-R₁-heteroaryl,

(14) - (CH₂)₀₋₄-CO-R₁-heterocycle wherein

R₁-heterocycle at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl, dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl, dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the R₁-heterocycle group is bonded by any atom of the parent R₁-heterocycle group substituted by hydrogen such that the new bond to the R₁-heterocycle group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(a) C₁-C₆ alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C₁-C₃ alkoxy,

(b) C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b}

(c) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b}

(d) halogen,

(e) C₁-C₆ alkoxy,

(f) -C₁-C₆ alkoxy optionally substituted with one, two, or three -F,

(g) $-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,(h) $-\text{OH}$,(i) $-\text{C}\equiv\text{N}$,(j) $(\text{CH}_2)_{0-4}-(\text{C}_3-\text{C}_7 \text{ cycloalkyl})$,

5 optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(k) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_4 \text{ alkyl})$,(l) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{1-a}\text{R}_{1-b}$,10 (m) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$,(n) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_6 \text{ alkyl})$, and(o) $=\text{O}$,(p) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{SO}_2-$ (q) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{C}(\text{O})-$ 15 (15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$ wherein

$\text{R}_{\text{N}-4}$ at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, pyrrolidinonyl, pyrrolyl, pyrazolyl, thienyl, pyridyl N-oxide, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl and pyrrolidinyl where each group is optionally substituted with 1, 2, 3, or 4 groups that are independently C_1-C_6 alkyl,

(16) $-(\text{CH}_2)_{0-4}-\text{CO}_2-\text{R}_{\text{N}-5}$ where

25 $\text{R}_{\text{N}-5}$ at each occurrence is independently selected from the group consisting of:

(a) C_1-C_6 alkyl,(b) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-aryl})$,(c) C_2-C_6 alkenyl,30 (d) C_2-C_6 alkynyl,(e) C_3-C_7 cycloalkyl, and(f) $-(\text{CH}_2)_{0-4}-(\text{R}_1\text{-heteroaryl})$,(17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$ (18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl})$,35 (19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,

- (20) - $(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3-\text{C}_7 \text{ cycloalkyl})$,
(21) - $(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}_2-\text{R}_{\text{N}-5}$,
(22) - $(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
(23) - $(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
(24) - $(\text{CH}_2)_{0-4}-\text{N}(-\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2}$,
(25) - $(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
(26) - $(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4}$,
(27) - $(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
(28) - $(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$ where R_{100} is
independently H or C_1-C_4 alkyl,
(29) - $(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
(30) - $(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
(31) - $(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
(32) - $(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
(33) - $(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
(34) - $(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})$ wherein the
alkyl group is optionally substituted with one, two, three,
four, or five substituents independently selected from the group
consisting of F, Cl, Br, and I,
(35) - $(\text{CH}_2)_{0-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
(36) C_2-C_6 alkenyl optionally substituted
with C_1-C_3 alkyl, halogen, -OH, -SH, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, or
 $-\text{NR}_{1-a}\text{R}_{1-b}$,
(37) C_2-C_6 alkynyl optionally substituted
with C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3
alkoxy, or $-\text{NR}_{1-a}\text{R}_{1-b}$, and
(38) - $(\text{CH}_2)_{0-4}-\text{N}(-\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$;
(IV) - $(\text{CR}_{\text{C}-x}\text{R}_{\text{C}-y})_{0-4}-\text{R}_{\text{C-heteroaryl}}$ wherein $\text{R}_{\text{C-heteroaryl}}$ at each
occurrence is independently selected from the group consisting
of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl,
indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl,
quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl,
pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl,
benzoisothiazolyl, benzimidazolyl, benzofuranyl, furanyl,
thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl,

tetrazolyl, oxazolopyridinyl, isothiazolyl, naphthyridinyl,
 cinnolinyl, carbazolyl, beta-carbolinyl, isochromanlyl, chromanlyl,
 tetrahydroisoquinolinyl, isoindolinyl,
 isobenzotetrahydrofuranyl, isobenzotetrahydrothienyl,
 5 isobenzothienyl, benzoxazolyl, pyridopyridinyl,
 benzotetrahydrofuranyl, benzotetrahydrothienyl, purinyl,
 benzodioxolyl, triazinyl, henoxazinyl, phenothiazinyl,
 pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
 dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
 10 dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl,
 coumarinyl, isocoumarinyl, chromonyl, chromanonyl,
 tetrahydroquinolinyl, dihydroquinolinyl,
 dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
 dihydroisocoumarinyl, isoindolinonyl,
 15 benzodioxanyl, benzoxazolinonyl, imidazopyrazolyl,
 quinazolinonyl, pyrazopyridyl, benzooxadiazolyl,
 dihydropyrimidinonyl, dihydrobenzofuranonyl,

where the R_C -heteroaryl group is bonded by any atom of the
 parent R_C -heteroaryl group substituted by hydrogen such that the new
 20 bond to the R_C -heteroaryl group replaces the hydrogen atom and its
 bond, where heteroaryl is optionally substituted 1, 2, 3, or 4
 groups that are independently:

(1) C_1 - C_6 alkyl, optionally substituted with 1, 2, or
 3 groups independently selected from the group consisting of C_1 -
 25 C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy,
 and -NR_{1-a}R_{1-b},

(2) -OH,

(3) -NO₂,

(4) -F, -Cl, -Br, -I,

30 (5) -CO-OH,

(6) -C \equiv N,

(V) C_2 - C_{10} alkenyl optionally substituted with one, two
 or three substituents independently selected from the group

consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, and -NR_{1-a}R_{1-b},

(VI) C₂-C₁₀ alkynyl optionally substituted with one, two or three substituents independently selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, and -NR_{1-a}R_{1-b},

(VII) -(C₁-C₆ alkyl)-O-(C₁-C₆ alkyl)-OH,

(VIII) -CH₂-NH-CH₂-CH(-O-CH₂-CH₃)₂,

(IX) -(CH₂)₀₋₆-C(=NR_{1-a})(NR_{1-a}R_{1-b});

10 where R_N is

(I) R_{N-1}-X_N- where X_N is -CO-, and where R_{N-1} is selected from the group consisting of:

(A) phenyl, which is optionally substituted with one, two or three of the following substituents which can be the same or different and are:

(1) C₁-C₆ alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

20 wherein R_{1-a} and R_{1-b} at each occurrence are independently H or C₁-C₆ alkyl,

(2) -OH,

(3) -NO₂,

(4) -F, -Cl, -Br, -I,

25 (5) -CO₂H,

(6) -C≡N,

(7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are the same or different and are selected from the group consisting of:

(a) -H,

30 (b) -C₁-C₈ alkyl optionally substituted with one substituent selected from the group consisting of:

(i) -OH,

(ii) -NH₂,

(iii) phenyl,

- (c) $-C_1-C_8$ alkyl optionally substituted with 1, 2, or 3 groups that are independently $-F$, $-Cl$, $-Br$, or $-I$,
- (d) $-C_3-C_8$ cycloalkyl,
- (e) $-(C_1-C_2 \text{ alkyl})-(C_3-C_8 \text{ cycloalkyl})$,
- 5 (f) $-(C_1-C_6 \text{ alkyl})-O-(C_1-C_3 \text{ alkyl})$,
- (g) $-C_2-C_6$ alkenyl,
- (h) $-C_2-C_6$ alkynyl,
- (i) $-C_1-C_6$ alkyl chain with one double bond and one triple bond,
- 10 (j) $-R_{1-aryl}$, wherein R_{1-aryl} at each occurrence is independently phenyl, naphthyl, indanyl, indenyl, dihydronaphthyl, or tetralinyl each of which is optionally substituted with 1, 2, 3, or 4 groups that are independently:
- (i) C_1-C_6 alkyl optionally
- 15 substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,
- (ii) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three
- 20 substituents independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,
- (iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,
- 25 (iv) $-F$, Cl , $-Br$ and $-I$,
- (v) $-C_1-C_6$ alkoxy optionally substituted with 1, 2, or 3 $-F$,
- (vi) $-NR_{N-2}R_{N-3}$,
- 30 (vii) $-OH$,
- (viii) $-C\equiv N$,
- (ix) C_3-C_7 cycloalkyl, optionally substituted with 1, 2, or 3 groups that are selected from the

group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(x) -CO-(C₁-C₄ alkyl),

(xi) -SO₂-NR_{1-a}R_{1-b},

5

(xii) -CO-NR_{1-a}R_{1-b}, or

(xiii) -SO₂-(C₁-C₄ alkyl),

(k) -R₁-heteroaryl, wherein R₁-heteroaryl at each occurrence is independently selected from the group consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, 10 indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, 15 triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranly, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, 20 pyridopyridinyl, benztetrahydrofuranly, benztetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl, dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl, dihydrobenzisothiazinyl, benzopyranly, benzothiopyranly, 25 coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-oxide, tetrahydroquinolinyl, dihydroquinolinyl, dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl, dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl, benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide, 30 pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide, indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide, quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide, thiazolyl N-oxide, indoliziny, indazolyl N-oxide, 35 benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-

oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is optionally substituted with 1, 2, 3, or 4 groups that are independently:

- (i) C_1-C_6 alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,
- (ii) C_2-C_6 alkenyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, or $-NR_{1-a}R_{1-b}$,
- (iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, or $-NR_{1-a}R_{1-b}$,
- (iv) -F, -Cl, -Br and -I,
- (v) $-C_1-C_6$ alkoxy optionally substituted with one, two, or three -F,
- (vi) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,
- (vii) -OH,
- (viii) $-C\equiv N$,
- (ix) $(CH_2)_{0-4}-C_3-C_7$ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,
- (x) $(CH_2)_{0-4}-CO-(C_1-C_6 \text{ alkyl})$,
- (xi) $(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,
- (xii) $(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,
- (xiii) $(CH_2)_{0-4}-SO_2-(C_1-C_6 \text{ alkyl})$,
- (xiv) $(CH_2)_{0-4}-N(R_{N-2})-SO_2-$, and
- (xv) $(CH_2)_{0-4}-N(R_{N-2})-C(O)-$,

(1) $-R_1$ -heterocycle, wherein

R_1 -heterocycle at each occurrence is independently selected from the group consisting of morpholinyl,

thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranlyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl, dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl, dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the R_1 -heterocycle group is bonded by any atom of the parent R_1 -heterocycle group substituted by hydrogen such that the new bond to the R_1 -heterocycle group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(a) C_1 - C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1 - C_3 alkyl, halogen, -OH, -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1 - C_3 alkoxy,

(b) C_2 - C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(c) C_2 - C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(d) halogen,

(e) C_1 - C_6 alkoxy,

(f) $-C_1$ - C_6 alkoxy optionally substituted with one, two, or three -F,

(g) $-NR_{N-2}R_{N-3}$,

(h) -OH,

(i) $-C\equiv N$,

- (j) $(\text{CH}_2)_{0-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
 optionally substituted with 1, 2, or 3 groups independently
 selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -
 CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},
- 5 (k) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_4 \text{ alkyl})$,
 (l) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{1-a}\text{R}_{1-b}$,
 (m) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$,
 (n) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_6 \text{ alkyl})$, and
 (o) =O,
- 10 (p) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{SO}_2-$
 (q) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{C}(\text{O})-$
- (8) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
 (9) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkenyl})$,
 (10) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkynyl})$,
 15 (11) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
 (12) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-aryl}$,
 (13) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl}$,
 (14) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle}$,
 (15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$ wherein R_{N-4} is selected from
 20 the group consisting of phenyl, morpholinyl, thiomorpholinyl,
 piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl,
 homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide,
 pyrrolinyl, thienyl, pyrazolyl, pyridyl N-oxide, oxazolyl,
 thiazolyl, imidazolyl, and pyrrolidinyl where each group is
 25 optionally substituted with one, two, three, or four groups that
 are independently C₁-C₆ alkyl,
- (16) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$ where R_{N-5} is selected
 from the group consisting of:
- (a) C₁-C₆ alkyl,
 30 (b) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-aryl})$,
 (c) C₂-C₆ alkenyl,
 (d) C₂-C₆ alkynyl,
 (e) $-(\text{CH}_2)_{0-2}-\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,
 (f) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heteroaryl})$, and
 35 (g) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heterocycle})$,

- (17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
- (18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl})$,
- (19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
- (20) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
- (21) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$,
- (22) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (23) $-(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (24) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2}$,
- (25) $-(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
- (26) $-(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4}$,
- (27) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
- (28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$ wherein
 R_{100} at each occurrence is independently -H
 or C_1-C_4 alkyl,
- (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
- (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
- (32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
- (33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
- (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl optionally substituted with one, two, three, four, or five of -F})$,
- (35) $\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,
- (36) $\text{C}_2-\text{C}_6 \text{ alkenyl optionally substituted with } \text{C}_1-\text{C}_3 \text{ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C}\equiv\text{N, -CF}_3, \text{C}_1-\text{C}_3 \text{ alkoxy, or -NR}_{1-a}\text{R}_{1-b}$,
- (37) $\text{C}_2-\text{C}_6 \text{ alkynyl optionally substituted with } \text{C}_1-\text{C}_3 \text{ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C}\equiv\text{N, -CF}_3, \text{C}_1-\text{C}_3 \text{ alkoxy, or -NR}_{1-a}\text{R}_{1-b}$,
- (38) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$,
- (39) $-(\text{CH}_2)_{1-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,

(B) $-\text{R}_{\text{N-heteroaryl}}$ where $\text{R}_{\text{N-heteroaryl}}$ is selected from the group consisting of pyridinyl, indolyl, indolinyl, isoindolyl, imidazolyl, isoxazolyl, oxazolyl, thiazolyl, indoliziny and isochromanyl,

where the $R_{N\text{-heteroaryl}}$ group is bonded by any atom of the parent $R_{N\text{-heteroaryl}}$ group substituted by hydrogen such that the new bond to the $R_{N\text{-heteroaryl}}$ group replaces the hydrogen atom and its bond, where heteroaryl is optionally substituted with one, two, three, or four of:

- (1) $C_1\text{-}C_6$ alkyl, optionally substituted with one, two or three substituents selected from the group consisting of $C_1\text{-}C_3$ alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, $C_1\text{-}C_3$ alkoxy, and $-NR_{1-a}R_{1-b}$,
- (2) $-OH$,
- (3) $-NO_2$,
- (4) $-F$, $-Cl$, $-Br$, $-I$,
- (5) $-CO_2H$,
- (6) $-C\equiv N$,
- (7) $-(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,
- (8) $-(CH_2)_{0-4}-CO-(C_1\text{-}C_{12} \text{ alkyl})$,
- (9) $-(CH_2)_{0-4}-CO-(C_2\text{-}C_{12} \text{ alkenyl})$,
- (10) $-(CH_2)_{0-4}-CO-(C_2\text{-}C_{12} \text{ alkynyl})$,
- (11) $-(CH_2)_{0-4}-CO-(C_3\text{-}C_8 \text{ cycloalkyl})$,
- (12) $-(CH_2)_{0-4}-CO-R_{1\text{-aryl}}$,
- (13) $-(CH_2)_{0-4}-CO-R_{1\text{-heteroaryl}}$,
- (14) $-(CH_2)_{0-4}-CO-R_{1\text{-heterocycle}}$,
- (15) $-(CH_2)_{0-4}-CO-R_{N-4}$,
- (16) $-(CH_2)_{0-4}-CO-O-R_{N-5}$,
- (17) $-(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,
- (18) $-(CH_2)_{0-4}-SO-(C_1\text{-}C_8 \text{ alkyl})$,
- (19) $-(CH_2)_{0-4}-SO_2-(C_1\text{-}C_{12} \text{ alkyl})$,
- (20) $-(CH_2)_{0-4}-SO_2-(C_3\text{-}C_8 \text{ cycloalkyl})$,
- (21) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO-O-R_{N-5}$,
- (22) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO-N(R_{N-5})_2$,
- (23) $-(CH_2)_{0-4}-N-CS-N(R_{N-5})_2$,
- (24) $-(CH_2)_{0-4}-N(-H \text{ or } R_{N-5})-CO-R_{N-2}$,
- (25) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,
- (26) $-(CH_2)_{0-4}-R_{N-4}$,
- (27) $-(CH_2)_{0-4}-O-CO-(C_1\text{-}C_6 \text{ alkyl})$,

- (28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$,
 (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
 5 (32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
 (33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
 (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6$ alkyl optionally substituted with one, two, three, four, or five of -F),
 (35) C_3-C_8 cycloalkyl,
 10 (36) C_2-C_6 alkenyl optionally substituted with C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, or $-\text{NR}_{1-a}\text{R}_{1-b}$,
 (37) C_2-C_6 alkynyl optionally substituted with C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy,
 15 or $-\text{NR}_{1-a}\text{R}_{1-b}$,
 (38) $-(\text{CH}_2)_{0-4}-\text{N}(-\text{H}$ or $\text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$,
 (39) $-(\text{CH}_2)_{1-4}-\text{C}_3-\text{C}_8$ cycloalkyl,
 (C) $\text{R}_{\text{N}}-\text{aryl}-\text{W}-\text{R}_{\text{N}}-\text{aryl}$,
 (D) $\text{R}_{\text{N}}-\text{aryl}-\text{W}-\text{R}_{\text{N}}-\text{heteroaryl}$,
 20 (E) $\text{R}_{\text{N}}-\text{aryl}-\text{W}-\text{R}_1-\text{heterocycle}$,
 (F) $\text{R}_{\text{N}}-\text{heteroaryl}-\text{W}-\text{R}_{\text{N}}-\text{aryl}$,
 (G) $\text{R}_{\text{N}}-\text{heteroaryl}-\text{W}-\text{R}_{\text{N}}-\text{heteroaryl}$,
 (H) $\text{R}_{\text{N}}-\text{heteroaryl}-\text{W}-\text{R}_{\text{N}-1}-\text{heterocycle}$,
 (I) $\text{R}_{\text{N}}-\text{heterocycle}-\text{W}-\text{R}_{\text{N}}-\text{aryl}$,
 25 (J) $\text{R}_{\text{N}}-\text{heterocycle}-\text{W}-\text{R}_{\text{N}}-\text{heteroaryl}$,
 (K) $\text{R}_{\text{N}}-\text{heterocycle}-\text{W}-\text{R}_{\text{N}-1}-\text{heterocycle}$,
 where W is
 (13) $-(\text{CH}_2)_{1-4}-$,
 (14) $-\text{O}-$,
 30 (15) $-\text{S}(\text{O})_{0-2}-$,
 (16) $-\text{N}(\text{R}_{\text{N}-5})-$,
 (17) $-\text{CO}-$; or
 (18) a bond;
 (II) $-\text{CO}-(\text{C}_1-\text{C}_6$ alkyl)-M-(C_1-C_6 alkyl), where M is S, SO or
 35 SO_2 , and wherein each alkyl is unsubstituted or substituted with

one, two, or three of substituents independently selected from the group consisting of:

(A) -NH-CO-(C₁-C₆ alkyl),

(B) -NH-CO-O-R_{N-8},

5 (C) -NR_{N-2}R_{N-3};

where R₁ is

-(CH₂)_{n₁}-phenyl, where n₁ is zero or one, and which is optionally substituted with one, two, three or four of the following substituents on the phenyl ring:

10 (A) C₁-C₆ alkyl optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

15 (B) C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

20 (C) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(D) -F, Cl, -Br or -I,

(F) -C₁-C₆ alkoxy optionally substituted with one, two or three of - F,

25 (G) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(H) -OH,

(I) -C≡N,

30 (J) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(K) -CO-(C₁-C₄ alkyl),

(L) -SO₂-NR_{1-a}R_{1-b},

(M) $-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$,

(N) $-\text{SO}_2-(\text{C}_1-\text{C}_4 \text{ alkyl})$; and

where R_2 is

(I) $-(\text{Z})-\text{C}_1-\text{C}_6 \text{ alkyl}$, where Z is a bond, $-\text{C}(\text{O})$, $-\text{CO}_2-$ or $-\text{SO}_2-$, wherein the alkyl group is optionally substituted with one, two or three substituents selected from the group consisting of $\text{C}_1-\text{C}_3 \text{ alkyl}$, $\text{C}_1-\text{C}_7 \text{ alkyl}$ (optionally substituted with $\text{C}_1-\text{C}_3 \text{ alkyl}$ and $\text{C}_1-\text{C}_3 \text{ alkoxy}$), $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, $-\text{NR}_{1-a}\text{R}_{1-b}$ where R_{1-a} and R_{1-b} are independently $-\text{H}$ or $\text{C}_1-\text{C}_6 \text{ alkyl}$, and $-\text{OC}=\text{O NR}_{1-a}\text{R}_{1-b}$,

(II) $-(\text{Z})-\text{CH}_2-\text{S}(\text{O})_{0-2}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,

(III) $-(\text{Z})-\text{CH}_2-\text{CH}_2-\text{S}(\text{O})_{0-2}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,

(IV) $-(\text{Z})-\text{C}_2-\text{C}_6 \text{ alkenyl}$ with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(V) $-(\text{Z})-\text{C}_2-\text{C}_6 \text{ alkynyl}$ with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(VI) $-(\text{Z})-(\text{CH}_2)_{n_1}-(\text{R}_{1-\text{aryl}})$, where Z is a bond, CO, CO_2 or SO_2 , where n_1 is zero or one and where $\text{R}_{1-\text{aryl}}$ is phenyl, 1-naphthyl, 2-naphthyl and indanyl, indenyl, dihydronaphthalenyl, or tetralinyl optionally substituted with one, two, three or four of the following substituents on the aryl ring:

(A) $\text{C}_1-\text{C}_6 \text{ alkyl}$ optionally substituted with one, two or three substituents selected from the group consisting of $\text{C}_1-\text{C}_3 \text{ alkyl}$, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(B) $\text{C}_2-\text{C}_6 \text{ alkenyl}$ with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(C) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

5 (D) -F, Cl, -Br or -I,

(F) -C₁-C₆ alkoxy optionally substituted with one, two or three of -F,

(G) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

10 (H) -OH,

(I) -C≡N,

(J) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -

15 NR_{1-a}R_{1-b},

(K) -CO-(C₁-C₄ alkyl),

(L) -SO₂-NR_{1-a}R_{1-b},

(M) -CO-NR_{1-a}R_{1-b},

(N) -SO₂-(C₁-C₄ alkyl),

20 (VII) -(Z)-(CH₂)_{n1}-(R_{1-heteroaryl}) where n₁ is as defined above and where R_{1-heteroaryl} is selected from the group consisting of:

pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, 25 quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, 30 isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl,

purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl,
pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl,
5 coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-
oxide, tetrahydroquinolinyl, dihydroquinolinyl,
dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl,
benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
10 pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,
indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide,
quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-
oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
thiazolyl N-oxide, indolizinyl N-oxide, indazolyl N-oxide,
15 benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide,
benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is bonded to $-(CH_2)_{n1}-$ by any ring atom
20 of the parent R_N -heteroaryl group substituted by hydrogen such that
the new bond to the R_1 -heteroaryl group replaces the hydrogen atom
and its bond, where heteroaryl is optionally substituted with
one, two, three or four of:

(1) C_1 - C_6 alkyl optionally substituted with one,
25 two or three substituents selected from the group consisting of
 C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH,
-SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(2) C_2 - C_6 alkenyl with one or two double bonds,
optionally substituted with one, two or three substituents
30 selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, -
 CF_3 , C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(3) C_2 - C_6 alkynyl with one or two triple bonds,
optionally substituted with one, two or three substituents

selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(4) -F, Cl, -Br or -I,

(6) -C₁-C₆ alkoxy optionally substituted with one, two, or three of -F,

(7) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(8) -OH,

(9) -C≡N,

(10) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(11) -CO-(C₁-C₄ alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

(13) -CO-NR_{1-a}R_{1-b}, or

(14) -SO₂-(C₁-C₄ alkyl), with the proviso that when n₁ is zero R_{1-heteroaryl} is not bonded to the carbon chain by nitrogen, or

(VIII) -(Z)-(CH₂)_{n1}-(R_{1-heterocycle}) where n₁ is as defined above and R_{1-heterocycle} is selected from the group consisting of:

morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl dihydropyrazinyl dihydropyridinyl dihydropyrimidinyl, dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, homothiomorpholinyl S-oxide,

where the R_{1-heterocycle} group is bonded by any atom of the parent R_{1-heterocycle} group substituted by hydrogen such that the new bond

to the R_1 -heterocycle group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with one, two, three or four:

(1) C_1 - C_6 alkyl optionally substituted with one, two or three substituents selected from the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

(2) C_2 - C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

(3) C_2 - C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

(4) -F, Cl, -Br, or -I,

(5) C_1 - C_6 alkoxy,

(6) - C_1 - C_6 alkoxy optionally substituted with one, two, or three -F,

(7) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(8) -OH,

(9) -C \equiv N,

(10) C_3 - C_7 cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

(11) -CO-(C_1 - C_4 alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

(13) -CO-NR_{1-a}R_{1-b},

(14) -SO₂-(C_1 - C_4 alkyl),

(15) =O, with the proviso that when n_1 is zero R_1 -heterocycle is not bonded to the carbon chain by nitrogen; and

where R₂₀ is H or C₁₋₆ alkyl or alkenyl.

The compounds of the invention, and pharmaceutically acceptable salts or esters thereof, are useful for treating humans who have Alzheimer's disease, for helping prevent or delay the onset of Alzheimer's disease, for treating patients with mild cognitive impairment (MCI) and preventing or delaying the onset of Alzheimer's disease in those who would progress from MCI to AD, for treating Down's syndrome, for treating humans who have Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, for treating cerebral amyloid angiopathy and preventing its potential consequences, i.e. single and recurrent lobar hemorrhages, for treating other degenerative dementias, including dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's disease, dementia associated with progressive supranuclear palsy, dementia associated with cortical basal degeneration, diffuse Lewy body type of Alzheimer's disease. It is preferred that the disease is Alzheimer's disease.

The compounds of the invention are also useful to inhibit beta-secretase and reduce or inhibit the formation of plaque.

When treating these diseases, compounds of the invention can either be used individually or together as is best for the patient.

With regard to these diseases the term "treating" means that compounds of the invention can be used in humans with existing disease. The compounds of the invention will not necessarily cure the patient who has the disease but will delay or slow the progression of the disease thereby giving the individual a more useful life span.

The term "preventing" means that that if the compounds of the invention are administered to those who do not now have the disease but who would normally get the disease or be at increased risk for the disease, they will not get the disease. In addition, "preventing" also includes delaying the development

of the disease in an individual who will ultimately get the disease or would be at risk for the disease. By delaying the onset of the disease, compounds of the invention have prevented the individual from getting the disease during the period in which the individual would normally have gotten the disease or reduce the rate of development of the disease or some of its effects but for the administration of compounds of the invention up to the time the individual ultimately gets the disease.

In treating or preventing the above diseases the compounds of the invention are administered in a therapeutically effective amount. The therapeutically effective amount will vary depending on the particular compound used and the route of administration as is known to those skilled in the art.

In treating a patient with any of the diagnosed above conditions a physician should begin administration of one or more of the compounds of the invention immediately and continue indefinitely.

In treating patients who do not at the have Alzheimer's disease, but who are believed to be at substantial risk for getting Alzheimer's disease in the future, the physician should start treatment when the patient first experiences early pre-Alzheimer's symptoms such as, memory or cognitive problems associated with aging. In addition, there are some patients who are at high risk because of having the genetic marker APOE4 which is predictive for Alzheimer's disease. In these situations, even though the patient does not have the disease, the administration of the compounds of the invention should be started before disease symptoms appear and treatment continued indefinitely to prevent or delay them from possibly getting the disease.

The compounds of the invention can be administered orally, parenterally (IV, IM, depo-IM, SQ and depo-SQ), sublingually, intranasally (inhalation), intrathecally, topically and rectally. The invention here is the compounds of the invention. There is nothing new about the routes of administration nor the

dosage forms. Dosage forms known to those skilled in the art are suitable for delivery of the compounds of the invention.

When administered orally, the compounds of the invention can be administered in usual dosage forms for oral administration as is well known to those skilled in the art. These dosage forms include the usual solid unit dosage forms of tablets and capsules as well as liquid dosage forms such as solutions, suspensions and elixirs. When the solid dosage forms are used, it is preferred that they be of the sustained release type so that the compounds of the invention need to be administered only once or twice daily.

The oral dosage forms are administered to the patient one thru four times daily. It is preferred that the compounds of the invention be administered either three or fewer time, more preferably once or twice daily. Hence, it is preferred that the compounds of the invention be administered in solid dosage form and further it is preferred that the solid dosage form be a sustained release form which permits once or twice daily dosing. It is preferred that whatever dosage form is used, that it be designed so as to protect the compounds of the invention from the acidic environment of the stomach. Enteric coated tablets are well known to those skilled in the art. In addition, capsules filled with small spheres each coated to protect from the acidic stomach, are also well known to those skilled in the art. When administered orally the therapeutically effective amount is from about 0.1 mg/day to about 1,000 mg/day. It is preferred that the oral dosage is from about 1 mg/day to about 100 mg/day. It is more preferred that the oral dosage is from about 5 mg/day to about 50 mg/day. It is understood that while a patient may be started on one dose, that dose may have to be varied over time as the patient's condition changes.

The compounds of the invention can be administered parenterally, for example, by IV, IM, depo-IM, SC, or depo-SC. When administered parenterally, a therapeutically effective amount of about 0.5 to about 100 mg/day, preferably from about 5

to about 50 mg daily should be delivered. When a depot formulation is used for injection once a month or once every two weeks, the dose should be about 0.5 mg/day to about 50 mg/day, or a monthly dose of from about 15 mg to about 1,500 mg. In part because of the forgetfulness of the patients with Alzheimer's disease, it is preferred that the parenteral dosage form be a depo formulation.

The compounds of the invention can be given sublingually. When given sublingually, the compounds of the invention should be given one thru four times daily in the same amount as for IM administration.

The compounds of the invention can be given intranasally. When given by this route of administration, the appropriate dosage forms are a nasal spray or dry powder as is known to those skilled in the art. The dosage of the compounds of the invention for intranasal administration is the same as for IM administration.

The compounds of the invention can be given intrathecally. When given by this route of administration the appropriate dosage form can be a parenteral dosage form as is known to those skilled in the art. The dosage of the compounds of the invention for intrathecal administration is the same as for IM administration.

The compounds of the invention can be administered topically. When given by this route, the appropriate dosage form is a cream, ointment, or patch. Because of the amount of the compounds of the invention to be administered, the patch is preferred. When administered topically, the dosage is from about 0.5 mg/day to about 200 mg/day. Because the amount that can be delivered by a patch is limited, two or more patches may be used. The number and size of the patch is not important, what is important is that a therapeutically effective amount of the compounds of the invention be delivered as is known to those skilled in the art. The compounds of the invention can be administered rectally by suppository as is known to those

skilled in the art. When administered by suppository, the therapeutically effective amount is from about 0.5 mg to about 500 mg.

5 The compounds of the invention can be administered by implants as is known to those skilled in the art. When administering a compound of the invention by implant, the therapeutically effective amount is the same as for depot administration.

10 The compounds of the invention are used in the same manner by the same routes of administration using the same pharmaceutical dosage forms and at the same dosing schedule for treating patients with MCI (mild cognitive impairment) and preventing or delaying the onset of Alzheimer's disease in those who would progress from MCI to AD, for treating Down's syndrome, 15 for treating humans who have Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, for treating cerebral amyloid angiopathy and preventing its potential consequences, i.e. single and recurrent lobar hemorrhages, for treating other degenerative dementias, including dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's 20 disease, dementia associated with progressive supranuclear palsy, dementia associated with cortical basal degeneration, diffuse Lewy body type of Alzheimer's disease.

25 The compounds of the invention can be used with each other or with other agents used to treat or prevent the conditions listed above. Such agents include gamma-secretase inhibitors, anti-amyloid vaccines and pharmaceutical agents such as donepezil hydrochloride (ARICEPT Tablets), tacrine hydrochloride 30 (COGNEX Capsules) or other acetylcholine esterase inhibitors and with direct or indirect neurotropic agents of the future.

In addition, the compounds of the invention can also be used with inhibitors of P-glycoprotein (P-gp). The use of P-gp inhibitors is known to those skilled in the art. See for 35 example, Cancer Research, 53, 4595-4602 (1993), Clin. Cancer

Res., 2, 7-12 (1996), Cancer Research, 56, 4171-4179 (1996), International Publications WO99/64001 and WO01/10387. The important thing is that the blood level of the P-gp inhibitor be such that it exerts its effect in inhibiting P-gp from decreasing brain blood levels of the compounds of the invention. To that end the P-gp inhibitor and the compounds of the invention can be administered at the same time, by the same or different route of administration, or at different times. The important thing is not the time of administration but having an effective blood level of the P-gp inhibitor.

Suitable P-gp inhibitors include cyclosporin A, verapamil, tamoxifen, quinidine, Vitamin E-TGPS, ritonavir, megestrol acetate, progesterone, rapamycin, 10,11-methanodibenzosuberane, phenothiazines, acridine derivatives such as GF120918, FK506, VX-710, LY335979, PSC-833, GF-102,918 and other steroids. It is to be understood that additional agents will be found that do the same function and are also considered to be useful.

The P-gp inhibitors can be administered orally, parenterally, (IV, IM, IM-depo, SQ, SQ-depo), topically, sublingually, rectally, intranasally, intrathecally and by implant.

The therapeutically effective amount of the P-gp inhibitors is from about 0.1 to about 300 mg/kg/day, preferably about 0.1 to about 150 mg/kg daily. It is understood that while a patient may be started on one dose, that dose may have to be varied over time as the patient's condition changes.

When administered orally, the P-gp inhibitors can be administered in usual dosage forms for oral administration as is known to those skilled in the art. These dosage forms include the usual solid unit dosage forms of tablets and capsules as well as liquid dosage forms such as solutions, suspensions and elixirs. When the solid dosage forms are used, it is preferred that they be of the sustained release type so that the P-gp inhibitors need to be administered only once or twice daily.

The oral dosage forms are administered to the patient one thru

four times daily. It is preferred that the P-gp inhibitors be administered either three or fewer times a day, more preferably once or twice daily. Hence, it is preferred that the P-gp inhibitors be administered in solid dosage form and further it is preferred that the solid dosage form be a sustained release form which permits once or twice daily dosing. It is preferred that what ever dosage form is used, that it be designed so as to protect the P-gp inhibitors from the acidic environment of the stomach. Enteric coated tablets are well known to those skilled in the art. In addition, capsules filled with small spheres each coated to protect From the acidic stomach, are also well known to those skilled in the art.

In addition, the P-gp inhibitors can be administered parenterally. When administered parenterally they can be administered IV, IM, depo-IM, SQ or depo-SQ. The P-gp inhibitors can be given sublingually. When given sublingually, the P-gp inhibitors should be given one thru four times daily in the same amount as for IM administration.

The P-gp inhibitors can be given intranasally. When given by this route of administration, the appropriate dosage forms are a nasal spray or dry powder as is known to those skilled in the art. The dosage of the P-gp inhibitors for intranasal administration is the same as for IM administration.

The P-gp inhibitors can be given intrathecally. When given by this route of administration the appropriate dosage form can be a parenteral dosage form as is known to those skilled in the art.

The P-gp inhibitors can be given topically. When given by this route of administration, the appropriate dosage form is a cream, ointment or patch. Because of the amount of the P-gp inhibitors needed to be administered the patch is preferred. However, the amount that can be delivered by a patch is limited. Therefore, two or more patches may be required. The number and size of the patch is not important, what is important is that a

therapeutically effective amount of the P-gp inhibitors be delivered as is known to those skilled in the art.

The P-gp inhibitors can be administered rectally by suppository as is known to those skilled in the art.

5 The P-gp inhibitors can be administered by implants as is known to those skilled in the art.

Route of administration and the dosage forms for administering the P-gp inhibitors are known in the art. Given a particular P-gp inhibitor, and a desired dosage form, one
10 skilled in the art would know how to prepare the appropriate dosage form for the P-gp inhibitor.

It should be apparent to one skilled in the art that the exact dosage and frequency of administration will depend on the particular compounds of the invention administered, the
15 particular condition being treated, the severity of the condition being treated, the age, weight, general physical condition of the particular patient, other medication the individual may be taking as is well known to those skilled in the art.

20 The compounds of the invention are also useful to inhibit beta-secretase and reduce or inhibit the formation of plaque.

Inhibition of APP Cleavage

The compounds of the invention inhibit cleavage of APP between Met595 and Asp596 numbered for the APP695 isoform, or a
25 mutant thereof, or at a corresponding site of a different isoform, such as APP751 or APP770, or a mutant thereof (sometimes referred to as the "beta secretase site". While not wishing to be bound by a particular theory, inhibition of beta-secretase activity is thought to inhibit production of beta
30 amyloid peptide (A-beta or Abeta). Inhibitory activity is demonstrated in one of a variety of inhibition assays, whereby cleavage of an APP substrate in the presence of A-beta-secretase enzyme is analyzed in the presence of the inhibitory compound, under conditions normally sufficient to result in cleavage at
35 the beta-secretase cleavage site. Reduction of APP cleavage at

the beta-secretase cleavage site compared with an untreated or inactive control is correlated with inhibitory activity. Assay systems that can be used to demonstrate efficacy of the compound inhibitors of the invention are known. Representative assay systems are described, for example, in U.S. Patents No. 5,942,400, 5,744,346, as well as in the examples below.

The enzymatic activity of beta-secretase and the production of Abeta can be analyzed in vitro or in vivo, using natural, mutated, and/or synthetic APP substrates, natural, mutated, and/or synthetic enzyme, and the test compound. The analysis may involve primary or secondary cells expressing native, mutant, and/or synthetic APP and enzyme, or may utilize transgenic animal models expressing the substrate and enzyme. Detection of enzymatic activity can be by analysis of one or more of the cleavage products, for example, by immunoassay, flurometric or chromogenic assay, HPLC, or other means of detection. Inhibitory compounds are determined as those having the ability to decrease the amount of beta-secretase cleavage product produced in comparison to a control, where beta-secretase mediated cleavage in the reaction system is observed and measured in the absence of inhibitory compounds.

Beta-secretase

Various forms of beta-secretase enzyme are known, and are available and useful for assay of enzyme activity and inhibition of enzyme activity. These include native, recombinant, and synthetic forms of the enzyme. Human beta-secretase is known as Beta Site APP Cleaving Enzyme (BACE), Asp2, and memapsin 2, and has been characterized, for example, in U.S. Patent 5,744,346 and published PCT patent applications WO98/22597, WO00/03819, WO01/23533, and WO00/17369, as well as in literature publications (Mol.Cell.Neurosci. 14:419-427 (1999); Science 286:735-741 (1999); Nature 402:533-537 (1999); Nature 40:537-540 (1999); and PNAS USA 97:1456-1460 (2000)). Synthetic forms of the enzyme have also been described (WO98/22597 and WO00/17369). Beta-secretase can be extracted and purified from human brain

tissue and can be produced in cells, for example mammalian cells expressing recombinant enzyme.

Preferred compounds of the invention are effective to inhibit 50% of beta-secretase enzymatic activity at a concentration of less than 50 micromolar, preferably at a concentration of 10 micromolar or less, more preferably 1 micromolar or less, and most preferably 10 nanomolar or less.

APP substrate

Assays that demonstrate inhibition of beta-secretase-mediated cleavage of APP can utilize any of the known forms of APP, including the 695 amino acid "normal" isotype described in Nature 325:733-6 (1987), the 770 amino acid isotype described in Nature 331:530-532 (1981), and variants such as the Swedish Mutation (KM670-1NL) (APP-SW), the London Mutation (V7176F), and others. See, for example U.S. Patent 5,766,846 and also Nature Genet. 1:233-234 (1992), for a review of known variant mutations. Additional useful substrates include the dibasic amino acid modification, APP-KK disclosed, for example, in WO 00/17369, fragments of APP, and synthetic peptides containing the beta-secretase cleavage site, wild type (WT) or mutated form, e.g., SW, as described, for example, in U.S. Patent 5,942,400 and WO00/03819.

The APP substrate contains the beta-secretase cleavage site of APP (KM-DA or NL-DA) for example, a complete APP peptide or variant, an APP fragment, a recombinant or synthetic APP, or a fusion peptide. Preferably, the fusion peptide includes the beta-secretase cleavage site fused to a peptide having a moiety useful for enzymatic assay, for example, having isolation and/or detection properties. A useful moiety may be an antigenic epitope for antibody binding, a label or other detection moiety, a binding substrate, and the like.

Antibodies

Products characteristic of APP cleavage can be measured by immunoassay using various antibodies, as described, for example, in Neuro. Lett. 249:21-4 (1999) and in U.S. Patent 5,612,486.

Useful antibodies to detect Abeta include, for example, the monoclonal antibody 6E10 (Senetek, St. Louis, MO) that specifically recognizes an epitope on amino acids 1-16 of the Abeta peptide; antibodies 162 and 164 (New York State Institute for Basic Research, Staten Island, NY) that are specific for human A-beta 1-40 and 1-42, respectively; and antibodies that recognize the junction region of beta-amyloid peptide, the site between residues 16 and 17, as described in U.S. Patent 5,593,846. Antibodies raised against a synthetic peptide of residues 591 to 596 of APP and SW192 antibody raised against 590-596 of the Swedish mutation are also useful in immunoassay of APP and its cleavage products, as described in U.S. Patents 5,604,102 and 5,721,130.

Assay Systems

Assays for determining APP cleavage at the beta-secretase cleavage site are well known in the art. Exemplary assays, are described, for example, in U.S. Patent 5,744,346 and 5,942,400, and described in the EXAMPLES below.

Cell free assays

Exemplary assays that can be used to demonstrate the inhibitory activity of the compounds of the invention are described, for example, in WO00/17369, WO 00/03819, and U.S. Patents 5,942,400 and 5,744,346. Such assays can be performed in cell-free incubations or in cellular incubations using cells expressing A-beta- secretase and an APP substrate having A-beta-secretase cleavage site.

An APP substrate containing the beat-secretase cleavage site of APP, for example, a complete APP or variant, an APP fragment, or a recombinant or synthetic APP substrate containing the amino acid sequence: KM-DA or NL-DA, is incubated in the presence of beta-secretase enzyme, a fragment thereof, or a synthetic or recombinant polypeptide variant having beta-secretase activity and effective to cleave the beta-secretase cleavage site of APP, under incubation conditions suitable for the cleavage activity of the enzyme. Suitable substrates

optionally include derivatives that may be fusion proteins or peptides that contain the substrate peptide and a modification useful to facilitate the purification or detection of the peptide or its beta-secretase cleavage products. Useful
5 modifications include the insertion of a known antigenic epitope for antibody binding; the linking of a label or detectable moiety, the linking of a binding substrate, and the like.

Suitable incubation conditions for a cell-free in vitro
10 assay include, for example: approximately 200 nanomolar to 10 micromolar substrate, approximately 10 to 200 picomolar enzyme, and approximately 0.1 nanomolar to 10 micromolar inhibitor compound, in aqueous solution, at an approximate pH of 4-7, at approximately 37°C, for a time period of approximately 10
15 minutes to 3 hours. These incubation conditions are exemplary only, and can be varied as required for the particular assay components and/or desired measurement system. Optimization of the incubation conditions for the particular assay components should account for the specific beta-secretase enzyme used and
20 its pH optimum, any additional enzymes and/or markers that might be used in the assay, and the like. Such optimization is routine and will not require undue experimentation.

One useful assay utilizes a fusion peptide having maltose binding protein (MBP) fused to the C-terminal 125 amino acids of
25 APP-SW. The MBP portion is captured on an assay substrate by anti-MBP capture antibody. Incubation of the captured fusion protein in the presence of beta-secretase results in cleavage of the substrate at the beta-secretase cleavage site. Analysis of the cleavage activity can be, for example, by immunoassay of
30 cleavage products. One such immunoassay detects a unique epitope exposed at the carboxy terminus of the cleaved fusion protein, for example, using the antibody SW192. This assay is described, for example, in U.S. Patent 5,942,400.

Cellular assay

Numerous cell-based assays can be used to analyze beta-secretase activity and/or processing of APP to release A-beta. Contact of an APP substrate with A-beta- secretase enzyme within the cell and in the presence or absence of a compound inhibitor of the invention can be used to demonstrate beta-secretase inhibitory activity of the compound. Preferably, assay in the presence of a useful inhibitory compound provides at least about 30%, most preferably at least about 50% inhibition of the enzymatic activity, as compared with a non-inhibited control.

In one embodiment, cells that naturally express beta-secretase are used. Alternatively, cells are modified to express a recombinant beta-secretase or synthetic variant enzyme as discussed above. The APP substrate may be added to the culture medium and is preferably expressed in the cells. Cells that naturally express APP, variant or mutant forms of APP, or cells transformed to express an isoform of APP, mutant or variant APP, recombinant or synthetic APP, APP fragment, or synthetic APP peptide or fusion protein containing the beta-secretase APP cleavage site can be used, provided that the expressed APP is permitted to contact the enzyme and enzymatic cleavage activity can be analyzed.

Human cell lines that normally process Abeta from APP provide a useful means to assay inhibitory activities of the compounds of the invention. Production and release of A-beta and/or other cleavage products into the culture medium can be measured, for example by immunoassay, such as Western blot or enzyme-linked immunoassay (EIA) such as by ELISA.

Cells expressing an APP substrate and an active beta-secretase can be incubated in the presence of a compound inhibitor to demonstrate inhibition of enzymatic activity as compared with a control. Activity of beta-secretase can be measured by analysis of one or more cleavage products of the APP substrate. For example, inhibition of beta-secretase activity against the substrate APP would be

expected to decrease release of specific beta-secretase induced APP cleavage products such as Abeta.

Although both neural and non-neural cells process and release A-beta, levels of endogenous beta-secretase activity are low and often difficult to detect by EIA. The use of cell types known to have enhanced beta-secretase activity, enhanced processing of APP to Abeta, and/or enhanced production of A-beta are therefore preferred. For example, transfection of cells with the Swedish Mutant form of APP (APP-SW); with APP-KK; or with APP-SW-KK provides cells having enhanced beta-secretase activity and producing amounts of A-beta that can be readily measured.

In such assays, for example, the cells expressing APP and beta-secretase are incubated in a culture medium under conditions suitable for beta-secretase enzymatic activity at its cleavage site on the APP substrate. On exposure of the cells to the compound inhibitor, the amount of Abeta released into the medium and/or the amount of CTF99 fragments of APP in the cell lysates is reduced as compared with the control. The cleavage products of APP can be analyzed, for example, by immune reactions with specific antibodies, as discussed above.

Preferred cells for analysis of beta-secretase activity include primary human neuronal cells, primary transgenic animal neuronal cells where the transgene is APP, and other cells such as those of a stable 293 cell line expressing APP, for example, APP-SW.

In vivo assays: animal models

Various animal models can be used to analyze beta-secretase activity and /or processing of APP to release Abeta, as described above. For example, transgenic animals expressing APP substrate and beta-secretase enzyme can be used to demonstrate inhibitory activity of the compounds of the invention. Certain transgenic animal models have been described, for example, in U.S. Patents 5,877,399, 5,612,486, 5,387,742, 5,720,936, 5,850,003, 5,877,015 and 5,811,633, and Nature 373:523 (1995)). Preferred are animals that exhibit characteristics associated

with the pathophysiology of AD. Administration of the compound inhibitors of the invention to the transgenic mice described herein provides an alternative method for demonstrating the inhibitory activity of the compounds. Administration of the compounds in a pharmaceutically effective carrier and via an administrative route that reaches the target tissue in an appropriate therapeutic amount is also preferred.

Inhibition of beta-secretase mediated cleavage of APP at the beta-secretase cleavage site and of Abeta release can be analyzed in these animals by measure of cleavage fragments in the animal's body fluids such as cerebral fluid or tissues. Analysis of brain tissues for Abeta deposits or plaques is preferred.

On contacting an APP substrate with A-beta-secretase enzyme in the presence of an inhibitory compound of the invention and under conditions sufficient to permit enzymatic mediated cleavage of APP and/or release of Abeta from the substrate, the compounds of the invention are effective to reduce beta-secretase-mediated cleavage of APP at the beta-secretase cleavage site and/or effective to reduce released amounts of Abeta. Where such contacting is the administration of the inhibitory compounds of the invention to an animal model, for example, as described above, the compounds are effective to reduce Abeta deposition in brain tissues of the animal, and to reduce the number and/or size of beta amyloid plaques. Where such administration is to a human subject, the compounds are effective to inhibit or slow the progression of disease characterized by enhanced amounts of Abeta to slow the progression of AD in the, and/or to prevent onset or development of AD in a patient at risk for the disease.

Unless defined otherwise, all scientific and technical terms used herein have the same meaning as commonly understood by one of skill in the art to which this invention belongs.

Definitions And Conventions

The definitions and explanations below are for the terms as used throughout this entire document including both the specification and the claims.

5 Definitions

Pharmaceutically acceptable refers to those properties and/or substances which are acceptable to the patient from a pharmacological/toxicological point of view and to the manufacturing pharmaceutical chemist from a physical/chemical
10 point of view regarding composition, formulation, stability, patient acceptance and bioavailability.

AD refers to Alzheimer's disease.

APP, amyloid precursor protein, is defined as any APP polypeptide, including APP variants, mutations, and isoforms,
15 for example, as disclosed in U.S. Patent 5,766,846.

A-beta (or Abeta), amyloid beta peptide, is defined as any peptide resulting from beta-secretase mediated cleavage of APP, including peptides of 39, 40, 41, 42, and 43 amino acids, and extending from the beta-secretase cleavage site to amino acids
20 39, 40, 41, 42, or 43.

Beta-secretase (beta-secretase, BACE1, Asp2, Memapsin 2) is an aspartyl protease that mediates cleavage of APP at the amino-terminal edge of Abeta. Human beta-secretase is described, for example, in WO00/17369.

25 DMSO refers to dimethyl sulfoxide.

All temperatures are in degrees Centigrade.

HPLC refers to high pressure liquid chromatography.

BOC refers to 1,1-dimethylethoxy carbonyl or *t*-butoxycarbonyl, -CO-O-C(CH₃)₃.

30 Protecting group generally refers to any suitable protecting groups, compatible with the synthetic routes for preparing the compounds herein. Generally, suitable protecting groups are those found in *Protective Groups in Organic Synthesis*, Greene, et. al., 2nd ed., John Wiley & Sons, 1991;
35 and 3rd ed., John Wiley & Sones, 1999 More specific protecting

groups are α -methyl benzyl, t-butoxycarbonyl, benzyloxycarbonyl, formyl, trityl, phthalimido, trichloroacetyl, chloroacetyl, bromoacetyl, iodoacetyl, 4-phenylbenzyloxycarbonyl, 2-methylbenzyloxycarbonyl, 4-ethoxybenzyloxycarbonyl, 4-fluorobenzyloxycarbonyl, 4-chlorobenzyloxycarbonyl, 3-chlorobenzyloxycarbonyl, 2-chlorobenzyloxycarbonyl, 2,4-dichlorobenzyloxycarbonyl, 4-bromobenzyloxycarbonyl, 3-bromobenzyloxycarbonyl, 4-nitrobenzyloxycarbonyl, 4-cyanobenzyloxycarbonyl, 2-(4-xenyl)isopropoxycarbonyl, 1,1-diphenyleth-1-yloxycarbonyl, 1,1-diphenylprop-1-yloxycarbonyl, 2-phenylprop-2-yloxycarbonyl, 2-(p-toluy)prop-2-yloxycarbonyl, cyclopentanyloxycarbonyl, 1-methylcycoopentanyloxycarbonyl, cyclohexanyloxycarbonyl, 1-methylcyclohexanyloxycabonyl, 2-methylcyclohexanyloxycarbonyl, 2-(4-toluylsulfonyl)ethoxycarbonyl, 2-(methy)lsulfonyl)ethoxycarbonyl, 2-(triphenylphosphino)ethoxycarbonyl, fluorenylmethoxycarbonyl, 2-(trimethylsilyl)ethoxycarbonyl, allyloxycarbonyl, 1-(trimethylsilylmethyl)prop-1-enyloxycarbonyl, 5-benzisoxalylmethoxycarbonyl, 4-acetoxybenzyloxycarbonyl, 2,2,2-trichloroethoxycarbonyl, 2-ethynyl-2-propoxycarbonyl, cyclopropylmethoxycarbonyl, 4-(decyloxy)benzyloxycarbonyl, isobronnyloxycarbonyl and 1-piperidyloxycarbonyl, 9-fluoroenylmethyl carbonate, $-\text{CH}=\text{CH}_2$ and phenyl-C(=N)-H.

Saline refers to an aqueous saturated sodium chloride solution.

Chromatography (column and flash chromatography) refers to purification/separation of compounds expressed as (support, eluent). It is understood that the appropriate fractions are pooled and concentrated to give the desired compound(s).

Pharmaceutically acceptable refers to those properties and/or substances that are acceptable to the patient from a pharmacological/toxicological point of view and to the manufacturing pharmaceutical chemist from a physical/chemical point of view regarding composition, formulation, stability, patient acceptance and bioavailability.

A therapeutically effective amount is defined as an amount effective to reduce or lessen at least one symptom of the disease being treated or to reduce or delay onset of one or more clinical markers or symptoms of the disease.

5 It should be noted that, as used in this specification and the appended claims, the singular forms "a," "an," and "the" include plural referents unless the content clearly dictates otherwise. Thus, for example, reference to a composition containing "a compound" includes a mixture of two or more
10 compounds. It should also be noted that the term "or" is generally employed in its sense including "and/or" unless the content clearly dictates otherwise.

Unless defined otherwise, all scientific and technical terms used herein have the same meaning as commonly understood
15 by one of skill in the art to which this invention belongs.

All patents and publications referred to herein are hereby incorporated by reference for all purposes.

Conventions for Formulas and Definitions of Variables

The chemical formulas representing various compounds or
20 molecular fragments in the specification and claims may contain variable substituents in addition to expressly defined structural features. These variable substituents are identified by a letter or a letter followed by a numerical subscript, for example, "Z₁" or "R_i" where "i" is an integer. These variable
25 substituents are either monovalent or bivalent, that is, they represent a group attached to the formula by one or two chemical bonds. For example, a group Z₁ would represent a bivalent variable if attached to the formula CH₃-C(=Z₁)H. Groups R_i and R_j would represent monovalent variable substituents if attached
30 to the formula CH₃-CH₂-C(R_i)(R_j)H₂. When chemical formulas are drawn in a linear fashion, such as those above, variable substituents contained in parentheses are bonded to the atom immediately to the left of the variable substituent enclosed in parentheses. When two or more consecutive variable substituents
35 are enclosed in parentheses, each of the consecutive variable

substituents is bonded to the immediately preceding atom to the left which is not enclosed in parentheses. Thus, in the formula above, both R_i and R_j are bonded to the preceding carbon atom. Also, for any molecule with an established system of carbon atom numbering, such as steroids, these carbon atoms are designated as C_i , where "i" is the integer corresponding to the carbon atom number. For example, C_6 represents the 6 position or carbon atom number in the steroid nucleus as traditionally designated by those skilled in the art of steroid chemistry. Likewise the term " R_6 " represents a variable substituent (either monovalent or bivalent) at the C_6 position.

Chemical formulas or portions thereof drawn in a linear fashion represent atoms in a linear chain. The symbol "-" in general represents a bond between two atoms in the chain. Thus $CH_3-O-CH_2-CH(R_i)-CH_3$ represents a 2-substituted-1-methoxypropane compound. In a similar fashion, the symbol "=" represents a double bond, e.g., $CH_2=C(R_i)-O-CH_3$, and the symbol " \equiv " represents a triple bond, e.g., $HC\equiv C-CH(R_i)-CH_2-CH_3$. Carbonyl groups are represented in either one of two ways: $-CO-$ or $-C(=O)-$, with the former being preferred for simplicity.

Chemical formulas of cyclic (ring) compounds or molecular fragments can be represented in a linear fashion. Thus, the compound 4-chloro-2-methylpyridine can be represented in linear fashion by $N^*=C(CH_3)-CH=CCl-CH=C^*H$ with the convention that the atoms marked with an asterisk (*) are bonded to each other resulting in the formation of a ring. Likewise, the cyclic molecular fragment, 4-(ethyl)-1-piperaziny1 can be represented by $-N^*-(CH_2)_2-N(C_2H_5)-CH_2-C^*H_2$.

A rigid cyclic (ring) structure for any compounds herein defines an orientation with respect to the plane of the ring for substituents attached to each carbon atom of the rigid cyclic compound. For saturated compounds which have two substituents attached to a carbon atom which is part of a cyclic system, $-C(X_1)(X_2)-$ the two substituents may be in either an axial or equatorial position relative to the ring and may change between

axial/equatorial. However, the position of the two substituents relative to the ring and each other remains fixed. While either substituent at times may lie in the plane of the ring (equatorial) rather than above or below the plane (axial), one substituent is always above the other. In chemical structural formulas depicting such compounds, a substituent (X_1) which is "below" another substituent (X_2) will be identified as being in the alpha configuration and is identified by a broken, dashed or dotted line attachment to the carbon atom, i.e., by the symbol
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"- - -" or "...". The corresponding substituent attached "above" (X_2) the other (X_1) is identified as being in the beta configuration and is indicated by an unbroken line attachment to the carbon atom. When a variable substituent is bivalent, the valences may be taken together or separately or both in the definition of the variable. For example, a variable R_i attached to a carbon atom as $-C(=R_i)-$ might be bivalent and be defined as oxo or keto (thus forming a carbonyl group $(-CO-)$ or as two separately attached monovalent variable substituents $\alpha-R_{i-j}$ and $\beta-R_{i-k}$. When a bivalent variable, R_i , is defined to consist of two monovalent variable substituents, the convention used to define the bivalent variable is of the form " $\alpha-R_{i-j}:\beta-R_{i-k}$ " or some variant thereof. In such a case both $\alpha-R_{i-j}$ and $\beta-R_{i-k}$ are attached to the carbon atom to give $-C(\alpha-R_{i-j})(\beta-R_{i-k})-$. For example, when the bivalent variable R_6 , $-C(=R_6)-$ is defined to consist of two monovalent variable substituents, the two monovalent variable substituents are $\alpha-R_{6-1}:\beta-R_{6-2}$, $\alpha-R_{6-9}:\beta-R_{6-10}$, etc, giving $-C(\alpha-R_{6-1})(\beta-R_{6-2})-$, $-C(\alpha-R_{6-9})(\beta-R_{6-10})-$, etc. Likewise, for the bivalent variable R_{11} , $-C(=R_{11})-$, two monovalent variable substituents are $\alpha-R_{11-1}:\beta-R_{11-2}$. For a ring substituent for which separate alpha and beta orientations do not exist (e.g. due to the presence of a carbon carbon double bond in the ring), and for a substituent bonded to a carbon atom which is not part of a ring the above convention is still used, but the alpha and beta designations are omitted.

Just as a bivalent variable may be defined as two separate monovalent variable substituents, two separate monovalent variable substituents may be defined to be taken together to form a bivalent variable. For example, in the formula $-C_1(R_i)H-C_2(R_j)H-$ (C_1 and C_2 define arbitrarily a first and second carbon atom, respectively) R_i and R_j may be defined to be taken together to form (1) a second bond between C_1 and C_2 or (2) a bivalent group such as oxa ($-O-$) and the formula thereby describes an epoxide. When R_i and R_j are taken together to form a more complex entity, such as the group $-X-Y-$, then the orientation of the entity is such that C_1 in the above formula is bonded to X and C_2 is bonded to Y . Thus, by convention the designation "... R_i and R_j are taken together to form $-CH_2-CH_2-O-CO-$..." means a lactone in which the carbonyl is bonded to C_2 . However, when designated "... R_j and R_i are taken together to form $-CO-O-CH_2-CH_2-$ the convention means a lactone in which the carbonyl is bonded to C_1 .

The carbon atom content of variable substituents is indicated in one of two ways. The first method uses a prefix to the entire name of the variable such as " C_1-C_4 ", where both "1" and "4" are integers representing the minimum and maximum number of carbon atoms in the variable. The prefix is separated from the variable by a space. For example, " C_1-C_4 alkyl" represents alkyl of 1 through 4 carbon atoms, (including isomeric forms thereof unless an express indication to the contrary is given). Whenever this single prefix is given, the prefix indicates the entire carbon atom content of the variable being defined. Thus C_2-C_4 alkoxy carbonyl describes a group $CH_3-(CH_2)_n-O-CO-$ where n is zero, one or two. By the second method the carbon atom content of only each portion of the definition is indicated separately by enclosing the " C_1-C_j " designation in parentheses and placing it immediately (no intervening space) before the portion of the definition being defined. By this optional convention (C_1-C_3) alkoxy carbonyl has the same meaning as C_2-C_4 alkoxy carbonyl because the " C_1-C_3 " refers only to the carbon

atom content of the alkoxy group. Similarly while both C₂-C₆ alkoxyalkyl and (C₁-C₃)alkoxy(C₁-C₃)alkyl define alkoxyalkyl groups containing from 2 to 6 carbon atoms, the two definitions differ since the former definition allows either the alkoxy or
5 alkyl portion alone to contain 4 or 5 carbon atoms while the latter definition limits either of these groups to 3 carbon atoms.

Modes of Preparation

Compounds of the invention can be prepared utilizing a
10 variety of known chemical transformations. In essence, the preparation of the compounds of formula I may be achieved using techniques and chemical processes analogous to those known in the art; the choice of the specific route being dependent upon the usual factors in pharmaceutical research institutions such
15 as availability and cost of starting materials, time and difficulties in separation and purification of intermediates and final compounds and such other factors well known and generally appreciated by those of ordinary skill in the art. In fact, there will generally be more than one process to prepare the
20 compounds of the invention. Those having skill in the art will recognize that the starting materials may be varied and additional steps employed to produce compounds encompassed by the present invention, as demonstrated by the following examples.

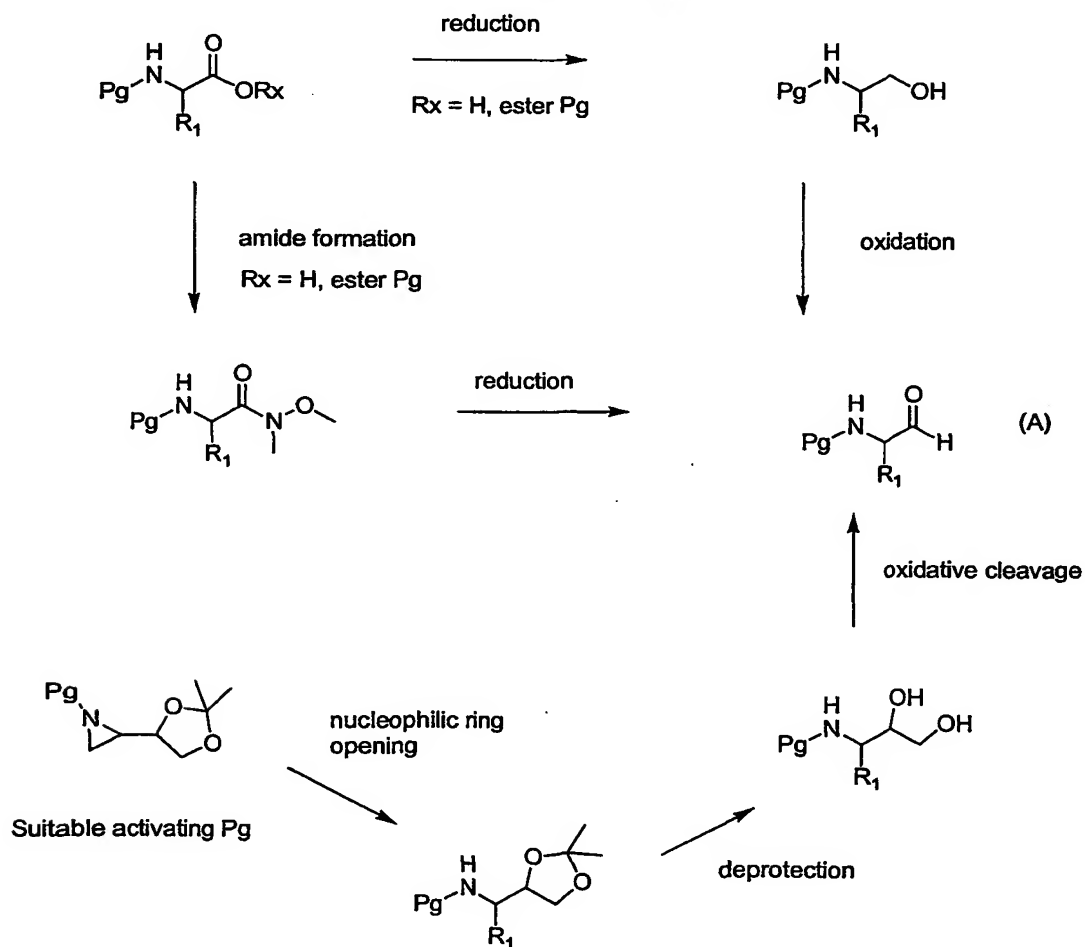
25 The starting materials and various intermediates may be obtained from commercial sources, prepared from commercially available organic compounds, or prepared using well known synthetic methods. In some cases, protection of reactive functionalities may be necessary to achieve some of the
30 transformations. In general, the need for such protecting groups as well as the conditions necessary to attach and remove such groups will be apparent to those skilled in the art of organic synthesis.

Representative synthetic methodologies suitable for
35 preparing the compounds of the invention are disclosed in:

Tetrahedron Letters, 39, 4925-4928 (1998); *Journal of Medicinal Chemistry*, 41, 3387-3401 (1998); *Journal of Medicinal Chemistry*, 39, 3203-3216 (1996); and/or *Journal of the Chemical Society Chemical Communications*, 1052-1053 (1993).

5 Examples of syntheses for preparing compounds of the invention are set forth below in Scheme 1, Scheme 2, Scheme 3 and Scheme 4. Specific exemplary descriptions of the routes depicted in Schemes 1 - 4 are found in the synthetic examples given for Intermediates A - D, general compound couplings E - G,
10 and examples 1 - 5 below.

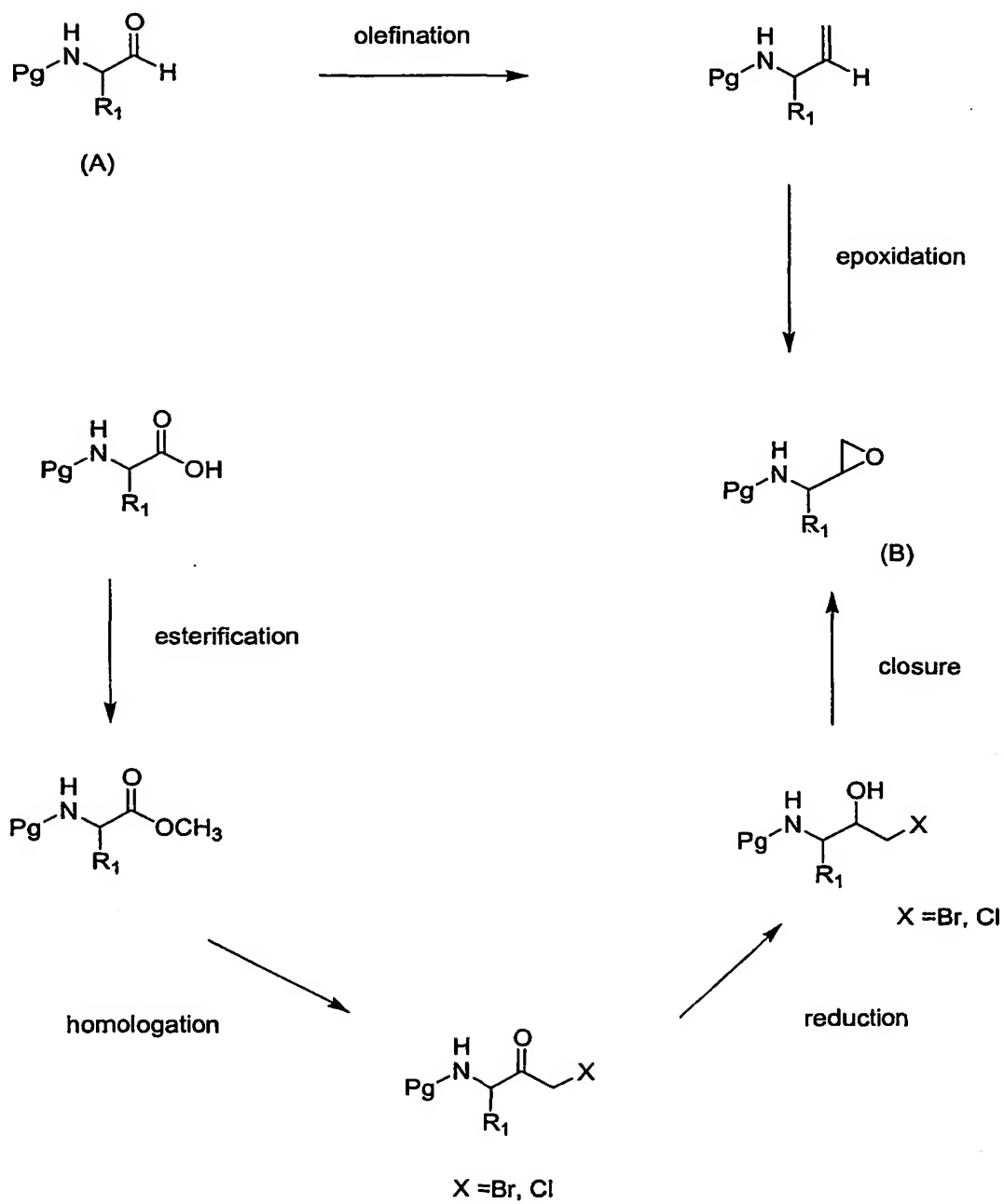
Scheme 1: Intermediate Synthesis



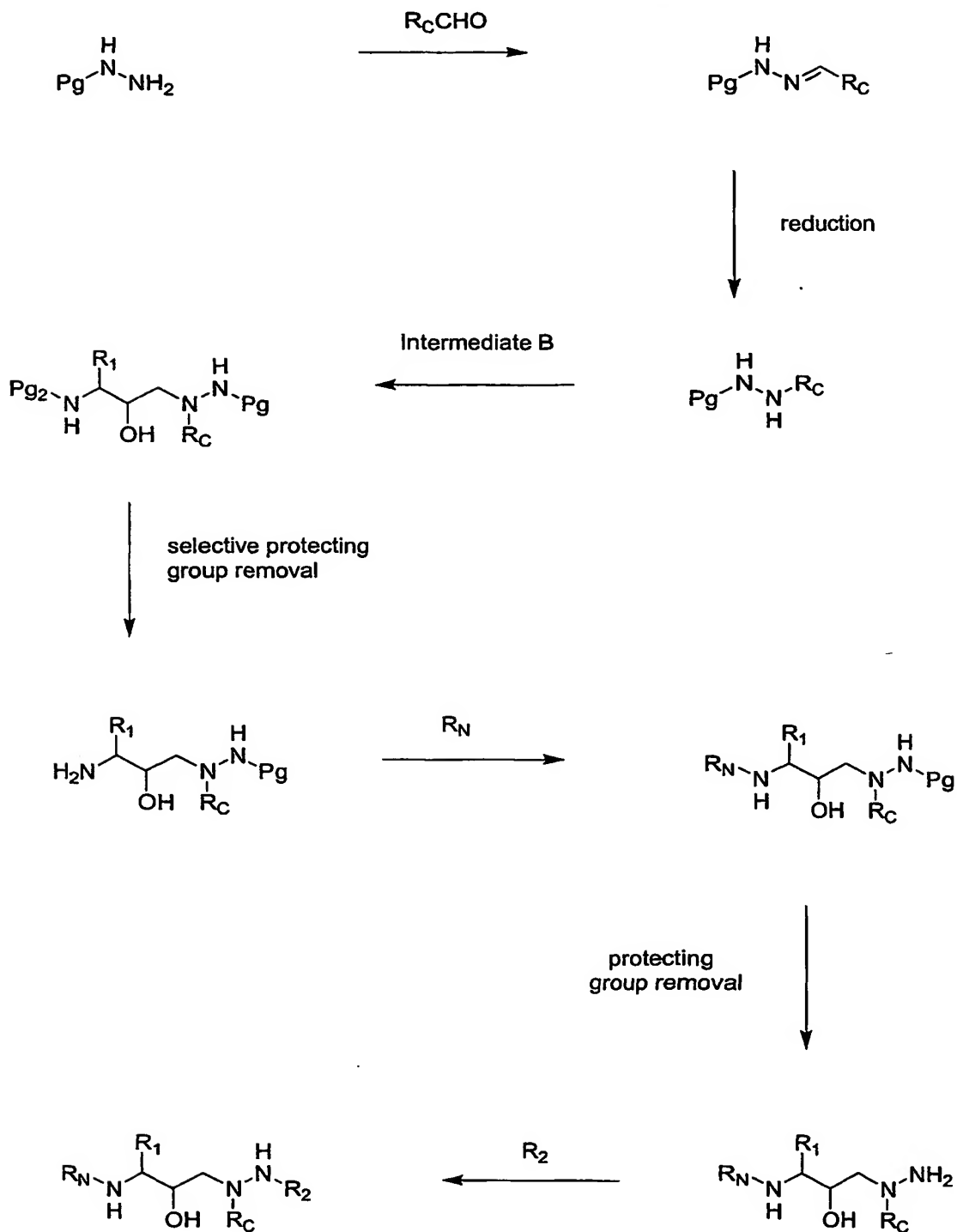
15 As used herein, Pg means protecting group, which is as defined herein; for exemplary synthesis of N-protected α -amino aldehydes see: *Chem. Rev.* 1989, 89, 149.

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Scheme 2: Intermediate Synthesis

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Scheme 3: General Compound Synthesis

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The invention is illustrated further by the following examples which are not to be construed as limiting the invention in scope or spirit to the specific procedures described in them.

5 Intermediate Syntheses

A. 2-Methanesulfonylamino-thiazole-4-carboxylic acid

2-Methanesulfonylamino-thiazole-4-carboxylic acid methyl ester: To a well stirred solution of methyl 2-amino-thiazole-4-carboxylate hydrobromide (*Justus Liebigs Ann. Chem.* 1951, 571, 44, *Heterocycles* 1997, 45, 1299) (13g, 51mmol) and TEA (40mL, 290mmol) in DCM (130mL) at rt was added methanesulfonyl chloride (11mL, 140mmol) dropwise. After stirring for 1h, the reaction mixture was filtered and the filtrate concentrated. The residue was re-suspended in ethyl acetate, filtered and chromatographed on silica gel (ethyl acetate/ethanol, 97:3) to provide 2-methanesulfonylamino-thiazole-4-carboxylic acid methyl ester (5.5g). MS (ESI-) for $C_6H_8N_2O_4S_2$ m/z 234.9 (M-H)⁻.

2-Methanesulfonylamino-thiazole-4-carboxylic acid: A mixture of 2-methanesulfonylamino-thiazole-4-carboxylic acid methyl ester (5.5g, 22mmol) in 1M NaOH (50mL) was stirred at rt for 1h. The reaction mixture was extracted with ethyl acetate, concentrated and chromatographed on silica gel (ethyl acetate/ethanol, 97:3) to provide 2-methanesulfonylamino-thiazole-4-carboxylic acid (3.7g) as a solid. MS (ESI-) for $C_5H_6N_2O_4S_2$ m/z 220.9 (M-H)⁻.

B. tert-Butyl 2-((2R,3S)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate

N'-Ethylidene-hydrazinecarboxylic acid tert-butyl ester: A solution of tert-butyl carbazate (50.0g, 0.38mole) and acetaldehyde (17.5g, 0.4mole) in ethanol (250mL) was refluxed for 72 h. The solution was concentrated in vacuo and dissolved in ethyl acetate. The solution was washed with water, dried (anhydrous sodium sulfate) and concentrated to afford N'-ethylidene-hydrazinecarboxylic acid tert-butyl ester (56g) as a

light yellow oil that solidified on standing. MS (ESI+) for $C_7H_{14}N_2O_2$ m/z 159.0 (M+H)⁺.

N'-Ethylhydrazinecarboxylic acid tert-butyl ester: A slurry of N'-ethylidene-hydrazinecarboxylic acid tert-butyl ester (5.0g, 32mmol), acetic acid (48 μ L) and PtO₂ (250mg) in ethanol (50mL) was shaken under a 40 psi pressure of H₂ in a Parr hydrogenator for 24h. The mixture was filtered through Celite to provide N'-ethylhydrazinecarboxylic acid tert-butyl ester (5g) as a clear oil. MS (ESI+) for $C_7H_{16}N_2O_2$ m/z 161.0 (M+H)⁺.

tert-Butyl 2-((2R,3S)-3-{[(benzyloxy) carbonyl] amino}-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate (*J. Med. Chem.* 1998, 41, 3387-3401): To a well stirred solution of [(1S)-1-(2S)-oxiranyl-2-phenylethyl]-carbamic acid phenylmethyl ester (6.07g, 20.4mmol) in isopropanol (100mL) was added N'-ethylhydrazinecarboxylic acid tert-butyl ester (3.93g, 24.5mmol) and the reaction refluxed for 67h. The reaction was concentrated and chromatographed on silica gel (elution with a gradient from hexane to dichloromethane to 1% methanol saturated with ammonia in dichloromethane) to give tert-butyl 2-((2R,3S)-3-{[(benzyloxy) carbonyl] amino}-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate (7.9g) as a white solid. MS (ESI+) for $C_{25}H_{35}N_3O_5$ m/z 458.3 (M+H)⁺.

tert-Butyl 2-((2R,3S)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate: A solution of tert-butyl 2-((2R,3S)-3-{[(benzyloxy) carbonyl] amino}-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate (3.11g, 6.8mmol) in methanol (50mL) was added to 10% Pd/C (0.32g) and stirred in a hydrogen atmosphere (@ ambient pressure) for 5h. The reaction was filtered through Celite and the filtrate concentrated under reduced pressure to yield tert-butyl 2-((2R,3S)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate (2.2g) as an oil. MS (ESI+) for $C_{17}H_{29}N_3O_3$ m/z 324.3 (M+H)⁺.

C. tert-Butyl 2-((2S,3S)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate

tert-Butyl 2-((2*S*,3*S*)-3-{[(benzyloxy)carbonyl]amino}-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate: To a well stirred solution of [(1*S*)-1-(2*R*)-oxiranyl-2-phenylethyl]-carbamic acid phenylmethyl ester (312mg, 1.0mmol) in isopropanol (3mL) was added *N'*-ethylhydrazinecarboxylic acid *tert*-butyl ester (202mg, 1.3mmol) and the reaction refluxed for 24h. The reaction was concentrated to give crude *tert*-butyl 2-((2*S*,3*S*)-3-{[(benzyloxy)carbonyl]amino

5]-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate (537mg). MS (ESI+) for C₂₅H₃₅N₃O₅ *m/z* 458.1 (M+H)⁺.

tert-Butyl 2-((2*S*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate: A solution of crude *tert*-butyl 2-((2*S*,3*S*)-3-{[(benzyloxy)carbonyl]amino}-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate (502mg, 1.1mmol) in MeOH (10mL) was added to 10% Pd/C (5.2mg) and stirred in a hydrogen atmosphere (@ ambient pressure) for 6h. The reaction was filtered through Celite and the filtrate concentrated under reduced pressure to yield crude *tert*-butyl 2-((2*S*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate (293mg). MS (ESI+) for C₁₇H₂₉N₃O₃ *m/z* 324.2 (M+H)⁺.

D. *tert*-Butyl 2-((2*S*,3*S*)-3-amino-2-hydroxy-4-(3,5-difluorophenyl)butyl)-2-ethylhydrazinecarboxylate

(*Z*)-2-Benzylloxycarbonylamino-3-phenyl-acrylate: A well stirred solution of 3,5-difluorobenzaldehyde (1.92g, 13.5mmol) and methyl benzyloxycarbonylamino-(dimethoxyphosphoryl)acetate (5.37g, 16.2mmol) in dry THF (20mL), under N₂, was cooled to 0°C and 1,1,3,3-tetramethylguanidine (1.38g, 16.2mmol) added via syringe. The reaction was allowed to warm to rt and stir for 6h, quenched with saturated ammonium chloride and extracted with ethyl acetate. After concentration, the residue was chromatographed on silica gel (elution with 20% ethyl acetate/heptane) to give methyl (*Z*)-2-benzyloxycarbonylamino-3-phenyl-acrylate (3.55g) as a white solid. MS (ESI+) for C₁₈H₁₅F₂NO₄ *m/z* 348.2 (M+H)⁺.

CBZ-L-3,5-Difluorophenylalanine Methyl Ester: A Parr Hastelloy bomb was charged with (Z)-2-benzyloxycarbonylamino-3-phenyl-acrylate (21g, 48mmol) in degassed methanol (315mL) and pressurized with H₂ @ 80 psi. After 1h, (S)-EtDuPHOSRh(COD)BF₄ (476mg, 1.5 mol%) in degassed methanol (15mL) was added and the reaction stirred at rt under 40 psi H₂ for 16h. The solution was filtered through Celite, concentrated and chromatographed on silica gel (elution with 20% ethyl acetate/heptane) to give 20g of CBZ-L-3,5-difluorophenylalanine methyl ester as a white solid. MS (ESI+) for C₁₈H₁₇F₂NO₄ m/z 350.1 (M+H)⁺.

[1S-(1-(3,5-Difluorobenzyl)-2-hydroxy-ethyl)]-carbamic acid benzyl ester: To a well stirred solution of L-3,5-difluorophenylalanine methyl ester (795mg, 2.3mmol) and lithium chloride (97mg, 2.3mmol) in THF/ethanol (10mL, 1: 1), under N₂, was added solid sodium borohydride (86mg, 2.3mmol) and the reaction stirred at rt overnight. The mixture was quenched with saturated ammonium chloride, extracted with ethyl acetate and the extracts combined, dried (anhydrous sodium sulfate) and concentrated. The residue was chromatographed on silica gel (elution with 20% ethyl acetate/heptane) to give [1S-(1-(3,5-difluorobenzyl)-2-hydroxy-ethyl)]-carbamic acid benzyl ester (660mg) as a white solid. MS (ESI+) for C₁₇H₁₇F₂NO₃ m/z 322.2 (M+H)⁺.

[1S-(1-(3,5-difluorobenzyl)-2-oxo-ethyl)]-carbamic acid benzyl ester: To a cold (-78 °C) well-stirred solution of oxalyl chloride (0.16 mL, 1.8mmol) in methylene chloride (20 mL) was added DMSO (0.13 mL, 1.8mmol) followed by the slow addition of [1S-(1-(3,5-difluoro-benzyl)-2-hydroxy-ethyl)]-carbamic acid benzyl ester (1.5mmol) as a solution in CH₂Cl₂ (10 mL). The reaction mixture was stirred at -78 °C for 10 minutes followed by the addition of Et₃N (0.63 mL, 4.5 mmol, 3.0 equiv). The mixture was stirred at -78 °C until no starting material was observed by TLC (2 h). The reaction mixture was then washed with 10% citric acid (aq) and dried over MgSO₄, filtered and condensed. The white solid was purified by silica chromatography

(20% to 50% EtOAc/ Heptane) to yield [1S-(1-(3,5-difluorobenzyl)-2-oxo-ethyl)]-carbamic acid benzyl ester (445 mg) as a white solid. MS (ESI+) for $C_{17}H_{15}F_2NO_3$ m/z 320 (M+H)⁺.

[1S-(1-(3,5-Difluorobenzyl)-2-propenyl)]-carbamic acid benzyl ester: To a slurry of diethyl ether (5mL) and Mg turnings (0.17g, 7.1mmol) was added $ClCH_2Si(CH_3)_3$ (0.97mL, 7.0 mmol) and the mixture was heated to reflux. After initiation (0.5h @ reflux), the reaction mixture was removed from heating and stirred for an additional 1 h, then cooled to rt. To the cooled solution was [1S-(1-(3,5-difluoro-benzyl)-2-oxo-ethyl)]-carbamic acid benzyl ester (0.45g, 1.4mmol) as a solution in distilled THF (10 mL). The reaction mixture was stirred at rt for 1 h, cooled to 0°C and $BF_3(OEt)_2$ (0.97mL, 7.0 mmol) added. The reaction mixture was further stirred at rt for 16 h, quenched with 10% NaOH (aq) (50mL) and stirring continued for 1h. The reaction mixture was extracted with CH_2Cl_2 , dried over $MgSO_4$, filtered and condensed to yield a clear oil. The oil was purified by silica chromatography (20% to 50% EtOAc/ heptane) to yield [1S-(1-(3,5-difluorobenzyl)-2-propenyl)]-carbamic acid benzyl ester (270 mg) as a white solid. MS (ESI+) for $C_{18}H_{18}F_2NO_2$ m/z 318 (M+H)⁺.

[(1S)-1-(2R)-oxiranyl-2-(3,5-difluorophenyl)ethyl]-carbamic acid phenylmethyl ester: To CH_2Cl_2 (20mL) was added [1S-(1-(3,5-difluoro-benzyl)-2-propenyl)]-carbamic acid benzyl ester (270mg, 0.85mmol) followed by M-CPBA (77%, 458mg, 1.0mmol). The reaction mixture was stirred at rt for 24h. The reaction was diluted with CH_2Cl_2 and then washed with 10% sodium thiosulfate (aq), dried over Na_2SO_4 , filtered and condensed. Preparative chromatography (5x50 cm Chiralcel OD, 30° C, 70 ml/min. 25% IPA/75% heptane) provided [(1S)-1-(2R)-oxiranyl-2-(3,5-difluorophenyl)ethyl]-carbamic acid phenylmethyl ester. MS (ESI+) for $C_{18}H_{18}F_2NO_3$ m/z 334.2 (M+H)⁺.

tert-Butyl 2-((2S,3S)-3-{[(benzyloxy)carbonyl]amino}-2-hydroxy-4-(3,5-difluorophenyl)butyl)-2-ethylhydrazinecarboxylate: To a well stirred solution of [(1S)-

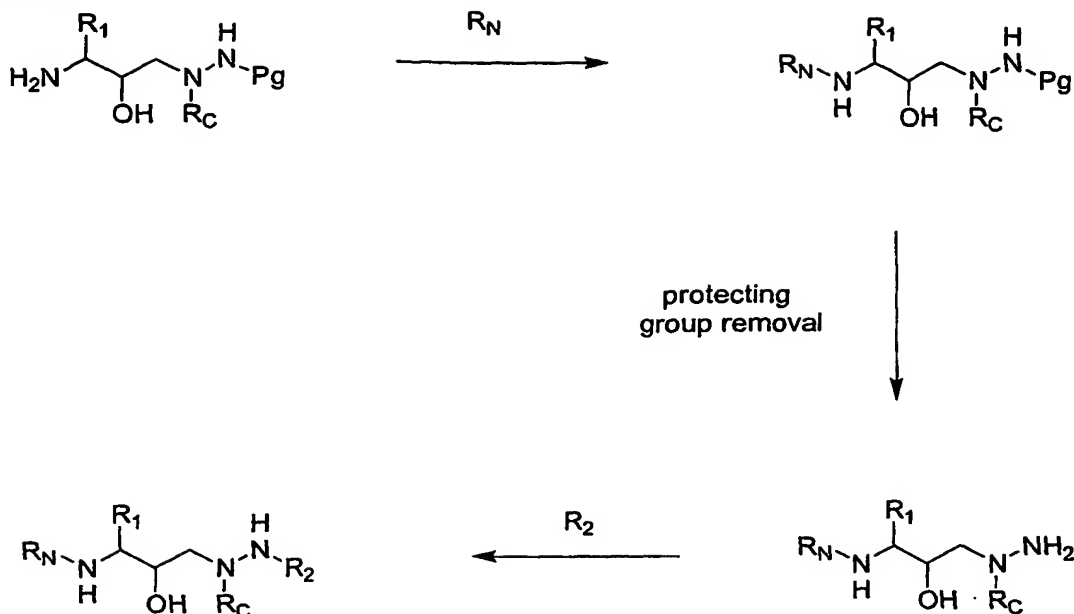
1- (2R) -oxiranyl-2- (3,5-difluorophenyl)ethyl]-carbamic acid
phenylmethyl ester (496mg, 1.5mmol) in isopropanol (4.25mL) was
added N'-ethylhydrazinecarboxylic acid tert-butyl ester (575mg,
3.6mmol) and the reaction refluxed for 48h. The reaction was
5 concentrated to provide crude tert-butyl 2-((2S,3S)-3-{
[(benzyloxy) carbonyl] amino}-2- hydroxy-4- (3,5-
difluorophenyl)butyl)-2-ethylhydrazine-carboxylate. MS (ESI+)
for C₂₅H₃₃F₂N₃O₅ m/z 516.2 (M+Na)⁺.

tert-Butyl 2-((2S,3S)-3-amino-2-hydroxy-4- (3,5-
10 difluorophenyl)butyl)-2- ethylhydrazinecarboxylate: A solution
of crude tert-butyl 2-((2S,3S)-3-{[(benzyloxy) carbonyl]-amino}-
2-hydroxy-4- (3,5-difluorophenyl)butyl)-2-
ethylhydrazinecarboxylate (906mg, 1.8mmol) in MeOH (15mL) was
added to 10% Pd/C (95mg) and stirred in a hydrogen atmosphere
15 (ambient pressure) for 4h. The catalyst was filtered off over
Celite and the filtrate was concentrated under reduced pressure
to yield crude tert-butyl 2-((2S,3S)-3-amino-2- hydroxy-4-
(3,5difluorophenyl)butyl)-2-ethylhydrazinecarboxylate. MS (ESI+)
for C₁₇H₂₇F₂N₃O₃ m/z 360.2 (M+H)⁺.

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Scheme 4: General Preparation Procedure

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E. N-terminal acid coupling (R_N)

5 Solutions of each acid (R_N) were prepared (1.08mmol) in DMF (3mL) (enough for 6 cartridges per acid). Solutions of PyBOP (3.74g, 7.2mmol) and HOBT (0.96g, 7.2mmol) in DMF (24mL) were prepared. To each cartridge on a Bohdan block was added acid solution (0.5mL, 0.18mmol) according to product layout (6

 10 cartridges of each of the 8 acids). To each cartridge was added the PyBOP/HOBT solution (0.5mL, 0.078g PYBOP, 0.15mmol/ 0.02g HOBT, 0.15mmol) and DIEA (0.052mL, 0.30mmol). The Bohdan block was agitated on a Bohdan shaker at 700-800 rpm for 1h. A solution of amine was prepared (2.30g, 7.2mmol) in 24mL of

 15 CH_2Cl_2 . To each cartridge was added 0.5mL of amine solution (0.48g, 0.15mmol). The Bohdan block was agitated on the Bohdan shaker at 700-800 rpm for 16h. The reaction mixtures were drained into 48 well Robbin's blocks. Each cartridge was rinsed with 0.5mL DMF into another 48 well Robbin's block. The reaction

 20 mixtures were concentrated in the Robbin's blocks under reduced pressure at 50 °C for 6h in a Jouan centrifugal evaporator. Products were carried on crude to next reaction.

F. Deprotection (*J. Org. Chem.* 1998, 63, 3471)

Add Dowex 50Wx2-400 ion-exchange resin (~230mg) to each clean cartridge on a Bohdan block using an Argo Scoop. The previous product was pipetted as a solution in 2mL MeOH into each cartridge. The Bohdan block was agitated on a Bohdan shaker at 50 °C at 700-800 rpm for 4h. Each cartridge was rinsed with CH₂Cl₂ (2x2.5mL) and MeOH (10x2.5mL). The products were eluted using 3.5M NH₃ in MeOH (2x3mL) into 2 separate 48 well Robbin's blocks. Reaction mixtures were concentrated in the Robbin's blocks under reduced pressure at 40 °C for 4h in a Jouan.

10 G. C-Terminal Acid Coupling (R₂)

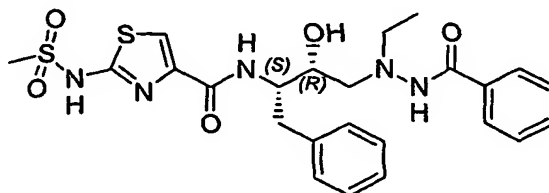
Solutions of each acid (R₂) were prepared (1.80mmol) in DMF (2mL) (enough for 8 cartridges per acid). A solution of HATU (4.13g, 10.8mmol) in DMF (12mL) was prepared. To each cartridge on a Bohdan block was added acid solution (0.25mL, 0.225mmol) according to product layout (8 cartridges of each of the 6 acids). To each cartridge was added 0.25mL of the HATU solution (0.086g, 0.225mmol) and DIEA (0.163mL, 0.94mmol). The Bohdan block was agitated on a Bohdan shaker at 700-800 rpm for 1h. Solution of the amines were prepared (0.15mmol/mL) in DMF. To each cartridge was added the amine solution (0.15mmol,) according to the product layout. The Bohdan block was agitated on the Bohdan shaker at 700-800 rpm for 16h. The reaction mixtures were drained into 48 well Robbin's blocks. Each cartridge was rinsed with 0.5mL DMF into another 48 well Robbin's block. The reaction mixtures were concentrated in the Robbin's blocks under reduced pressure at 50 °C for 6h in a Jouan. Products were dissolved in 2mL MeOH and each transferred to a Varian Mega BE-SCX (2gm, 12mL) cartridge that had been presoaked with MeOH (5mL). After products were loaded onto SCX cartridges by gravity, each cartridge was washed with MeOH (9x5mL) using vacuum filtration. Final products were eluted off by gravity using 3.5M NH₃ in MeOH (3x3mL) into pre-tared vials.

Examples

Example 1. N-[(1S,2R)-3-(2-benzoyl-1-ethylhydrazino)-1-benzyl-2-hydroxypropyl]-2-

[(methylsulfonyl)amino]-1,3-thiazole-4-carboxamide: Synthesized as described above from *tert*-butyl 2-((2*R*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate, benzoic acid and 2-methanesulfonylamino-thiazole-4-carboxylic acid. MS (ESI+) for

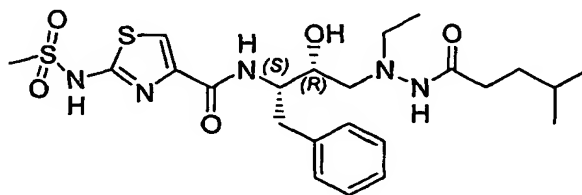
5 $C_{24}H_{29}N_5O_5S_2$ m/z 532.7 (M+H)⁺.



10

Example 2. N-{(1*S*,2*R*)-1-benzyl-3-[1-ethyl-2-(4-methylpentanoyl)hydrazino]-2-hydroxypropyl}-2-[(methylsulfonyl)amino]-1,3-thiazole-4-carboxamide: Synthesized as described above from *tert*-butyl 2-((2*R*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate, 4-methylpentanoic acid and 2-methanesulfonylamino-thiazole-4-carboxylic acid.. MS (ESI+) for

15 $C_{23}H_{35}N_5O_5S_2$ m/z 526.7 (M+H)⁺.

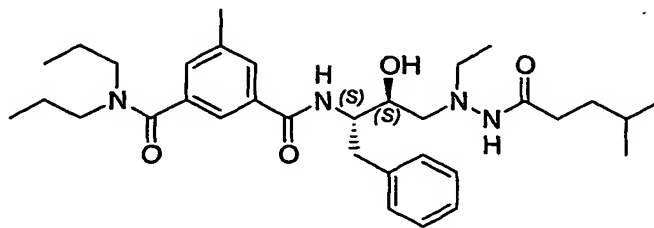


20

Example 3. N¹-{(1*S*,2*S*)-1-benzyl-3-[1-ethyl-2-(4-methylpentanoyl)hydrazino]-2-hydroxypropyl}-5-methyl-N³,N³-dipropylisophthalamide: Synthesized as described above from *tert*-butyl 2-((2*S*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl)-2-ethylhydrazinecarboxylate, 4-

25

methylpentanoic acid and 5-methyl-N,N-dipropyl-isophthalamide acid (WO 02/02512). MS (ESI+) for $C_{33}H_{50}N_4O_4$ m/z 567.3 $(M+H)^+$.



5

Example 4. N^1 -{ (1S,2S) -1-(3,5-difluorobenzyl) -3-[1-ethyl-2-(4-methylpentanoyl)hydrazino] -2-

10 hydroxypropyl} -5-methyl- N^3,N^3 -dipropylisophthalamide:

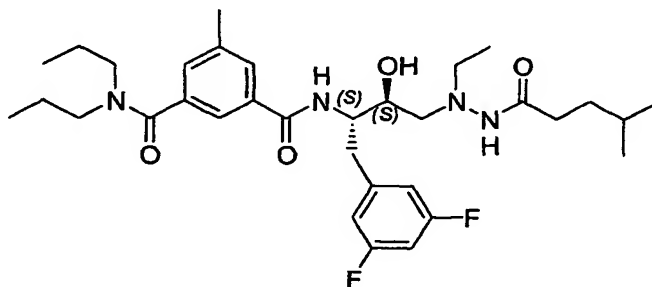
Synthesized as described above from tert-butyl 2-((2S,3S)-3-amino-2-

hydroxy-4-(3,5difluorophenyl)butyl)-2-

ethylhydrazinecarboxylate, 3-methylpentanoic acid and 5-methyl-

N,N-dipropyl-isophthalamide acid. MS (ESI+) for $C_{33}H_{48}F_2N_4O_4$ m/z

15 603.3 $(M+H)^+$.



20 Example 5. N^1 -{ (1S,2S) -1-(3,5-difluorobenzyl) -3-[1-ethyl-2-(4-methylbutanoyl)hydrazino] -2-

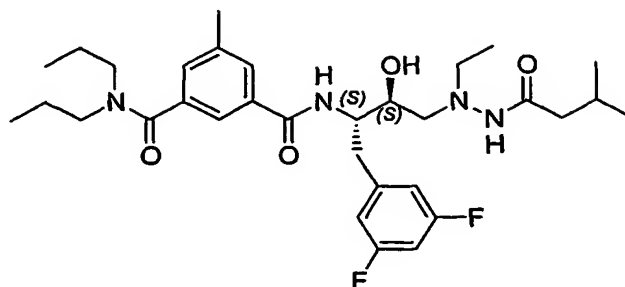
hydroxypropyl} -5-methyl- N^3,N^3 -dipropylisophthalamide:

Synthesized as described above from tert-butyl 2-((2S,3S)-3-amino-2-

hydroxy-4-(3,5difluorophenyl)butyl)-2-

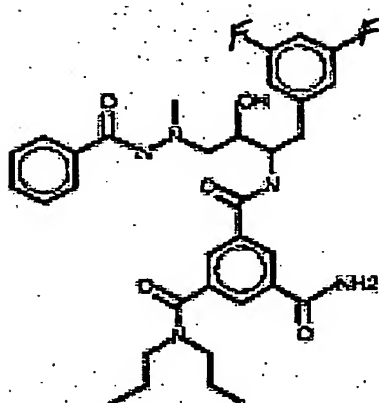
25 ethylhydrazinecarboxylate, 3-methylbutanoic acid and 5-methyl-

N,N-dipropyl-isophthalamide. MS (ESI+) for $C_{32}H_{46}F_2N_4O_4$ m/z 589.3 (M+H)⁺.

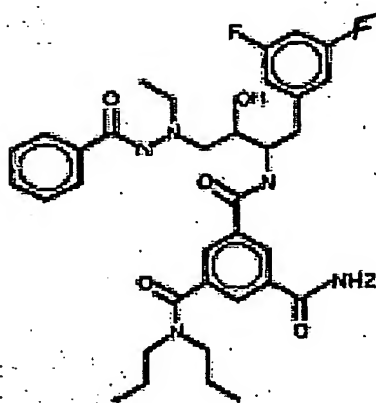


Example 6. The following compounds, as shown in Table 1 below, are prepared essentially according to the procedures outlined in Schemes 1-4 and according to examples 1-5:

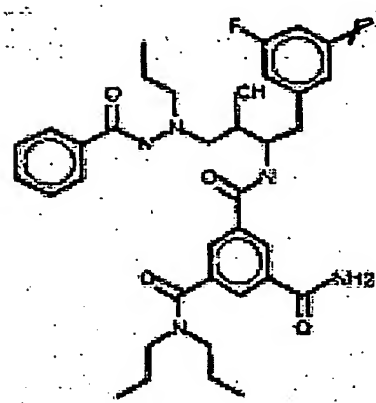
Table 1



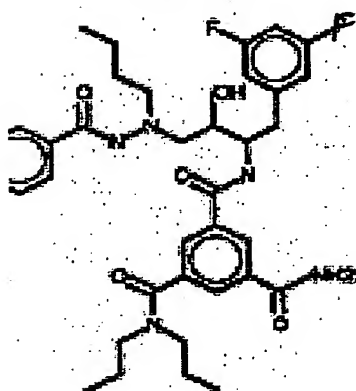
1A1 1,1,3,1



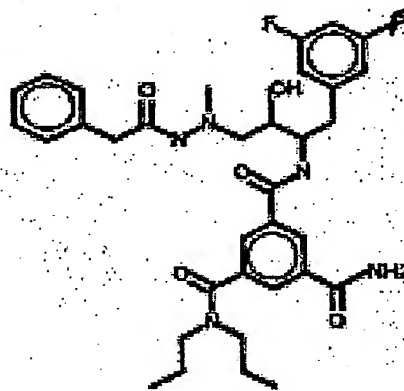
1A2 1,1,1,2



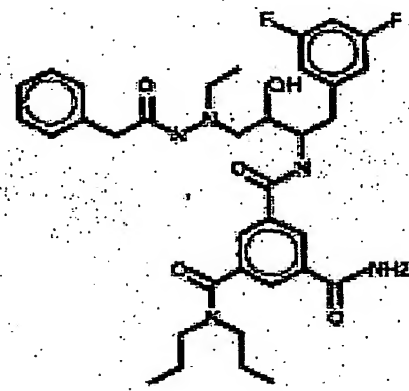
1A3 1,1,1,2



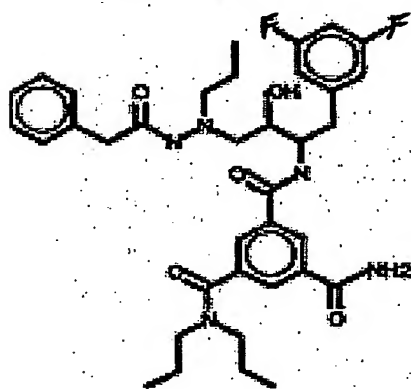
1A4 1,1,3,4



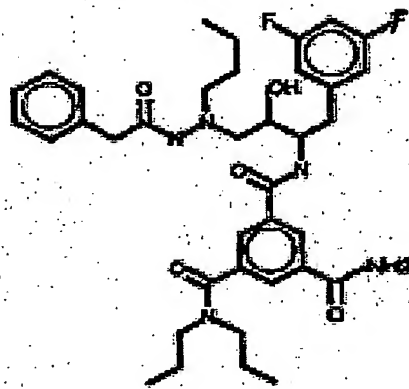
1A5 1,1,2,1



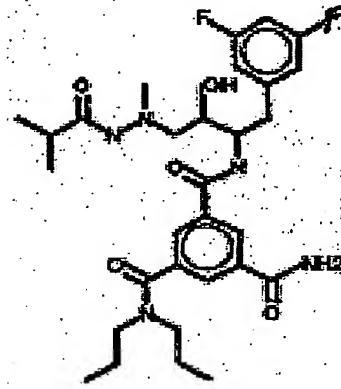
1A6 1,1,2,2



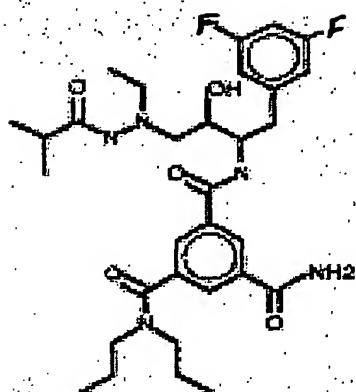
1A7 1,1,2,3



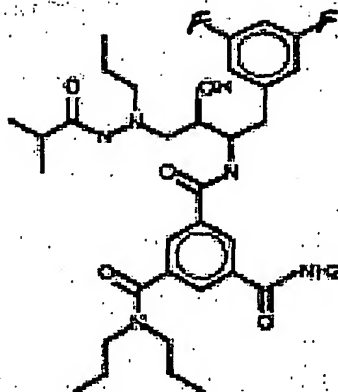
1A8 1,1,2,4



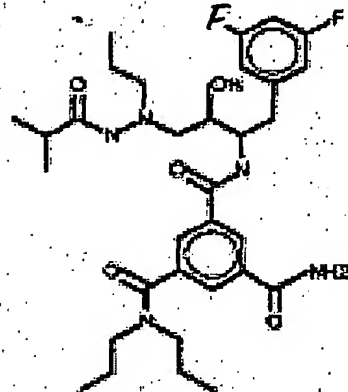
1A9 1,1,3,1



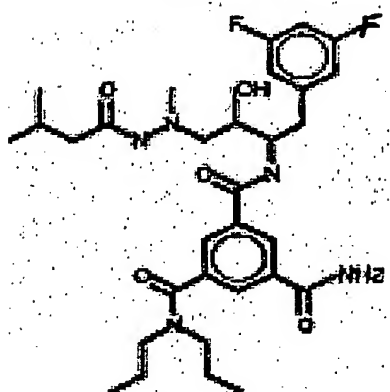
1A10 1,1,3,2



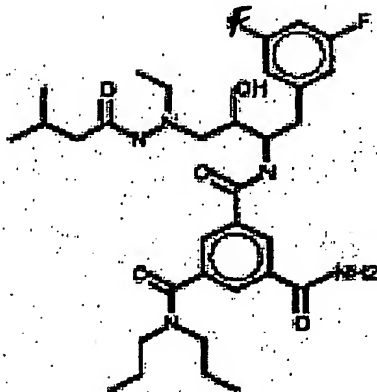
1A11 1,1,3,3



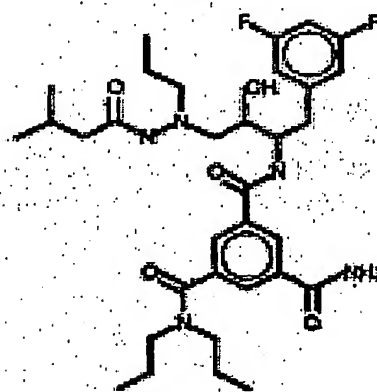
1A12 1,1,3,4



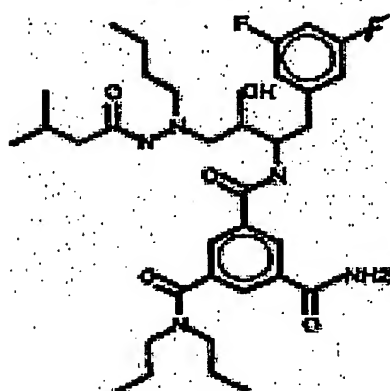
1B1 1,1,4,1



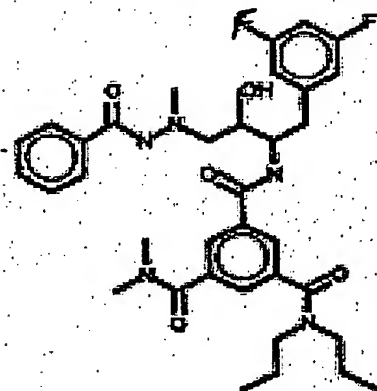
1B2 1,1,4,2



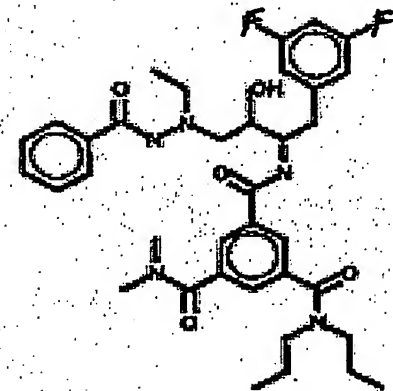
1B3 1,1,4,3



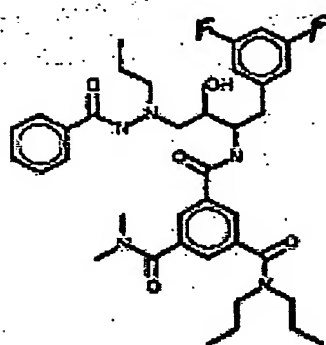
1B4 1,1,4,4



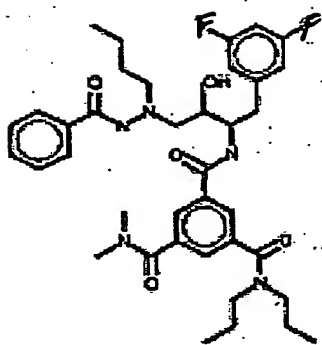
1B5 1,2,1,1



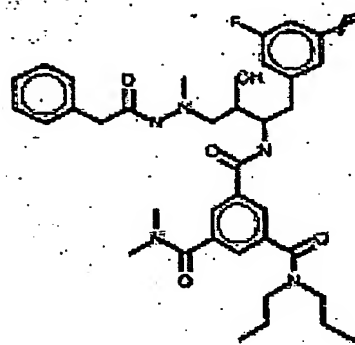
1B6 1,2,1,2



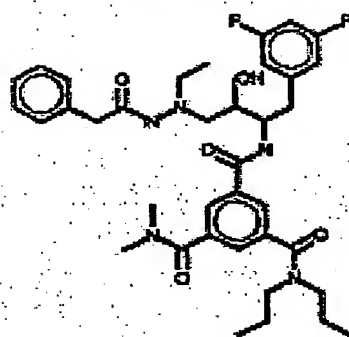
1:87 1,2,1,3



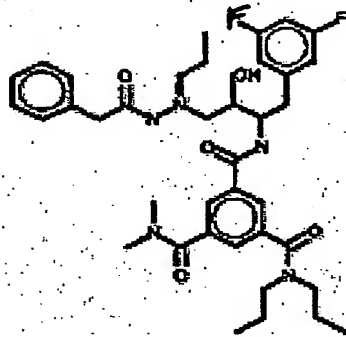
1:88 1,2,1,4



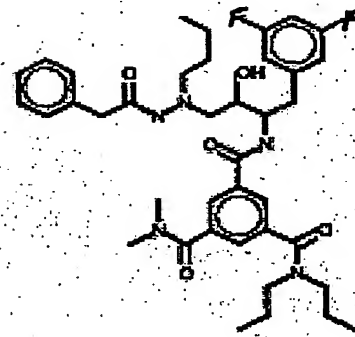
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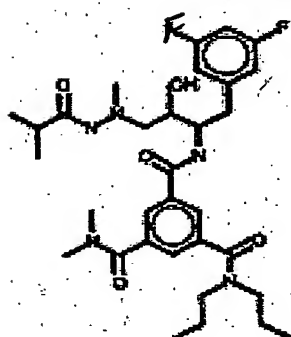
1:810 1,2,2,2



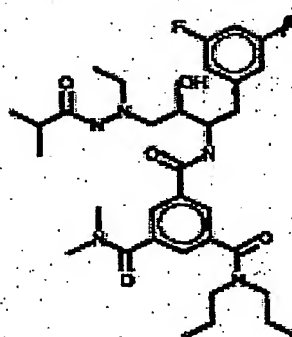
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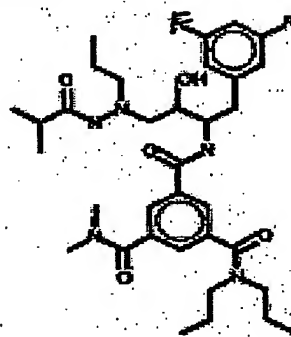
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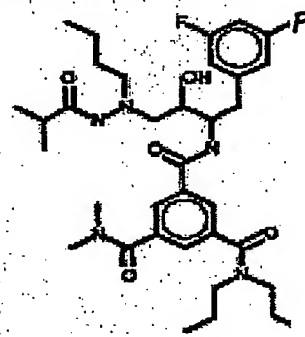
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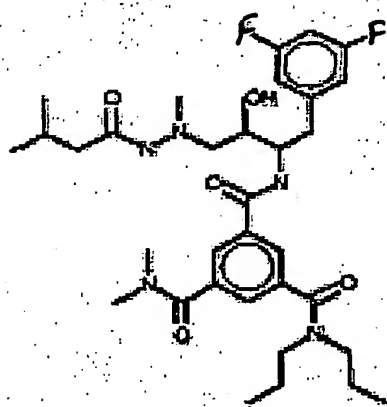
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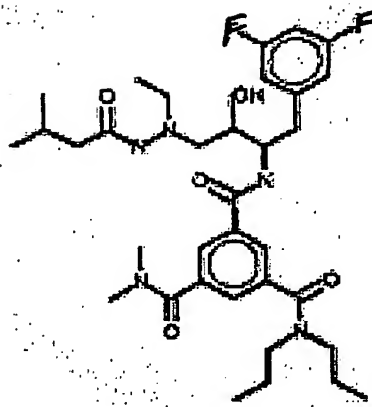
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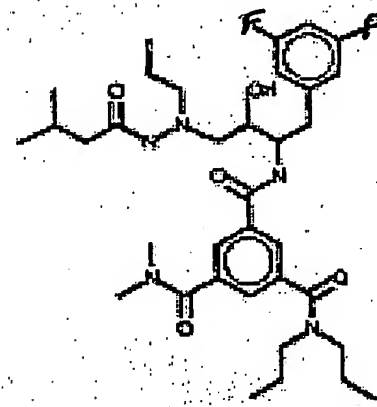
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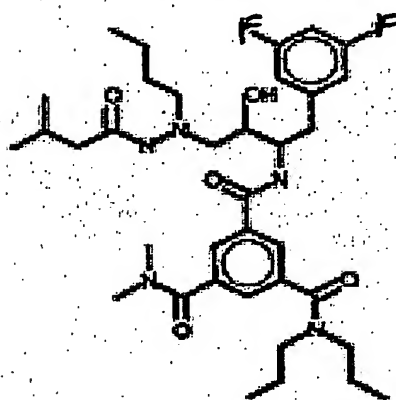
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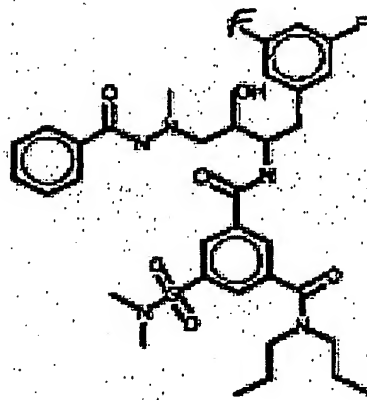
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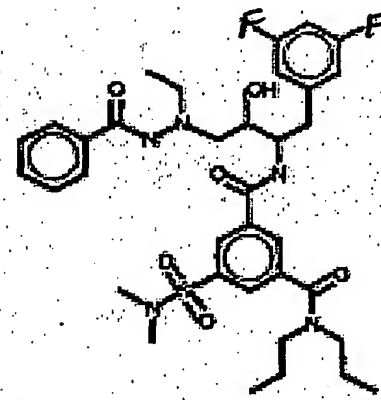
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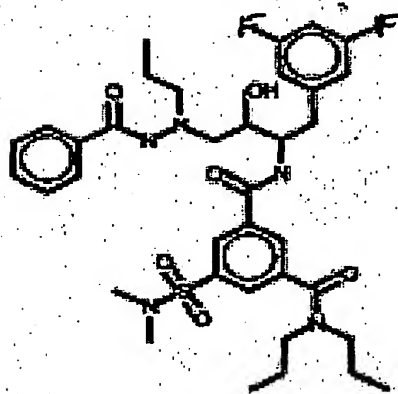
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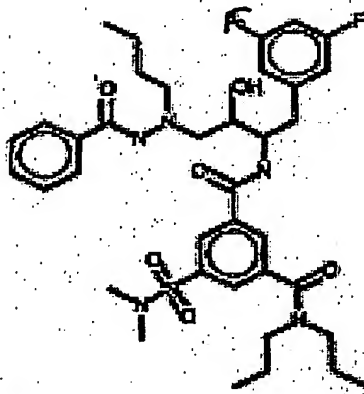
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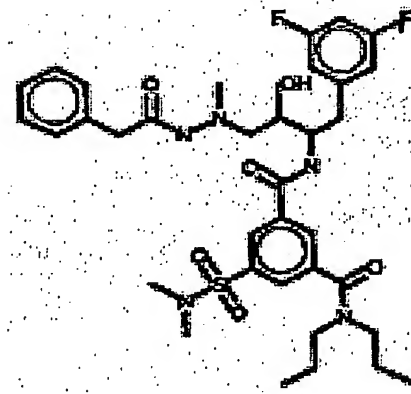
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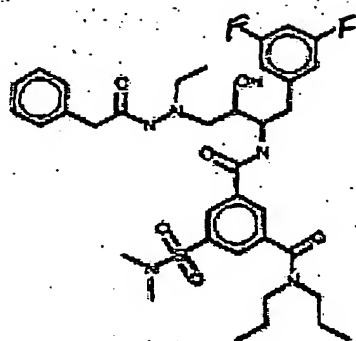
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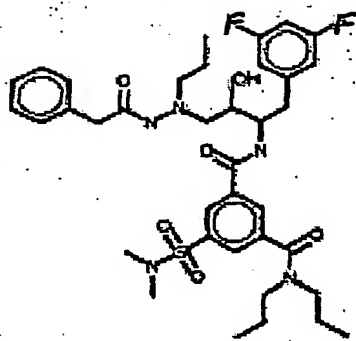
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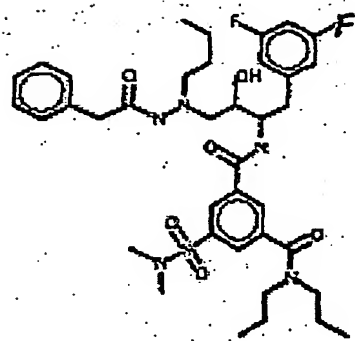
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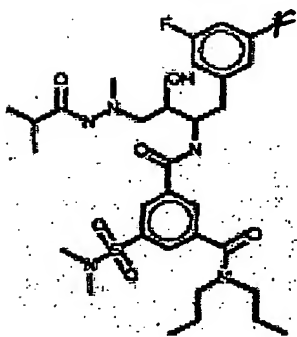
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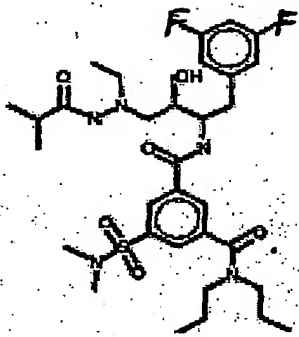
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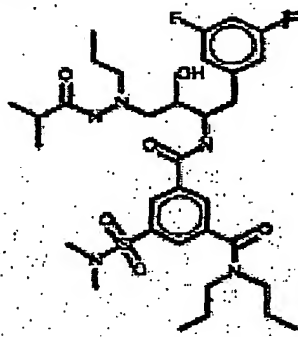
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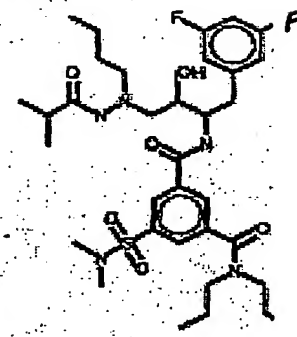
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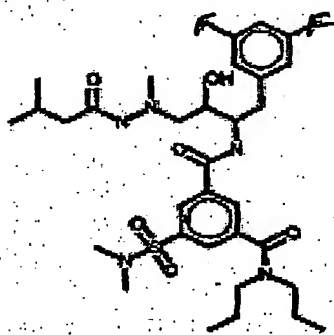
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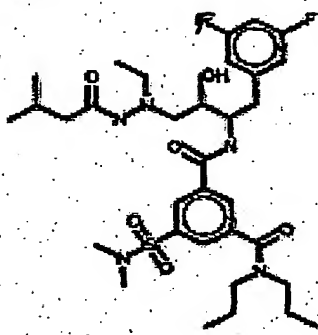
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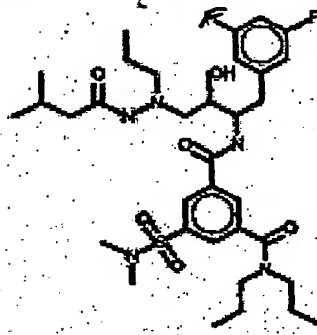
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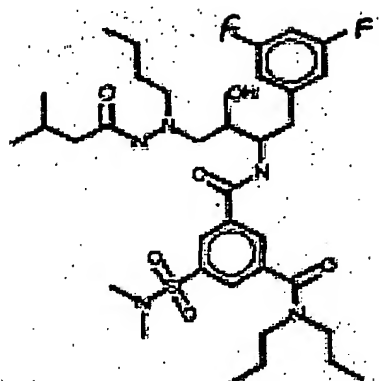
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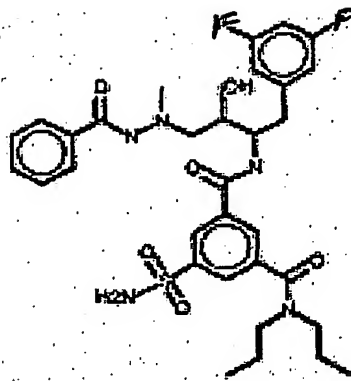
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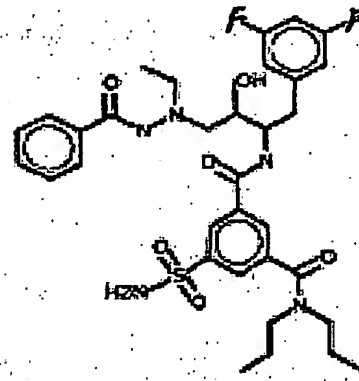
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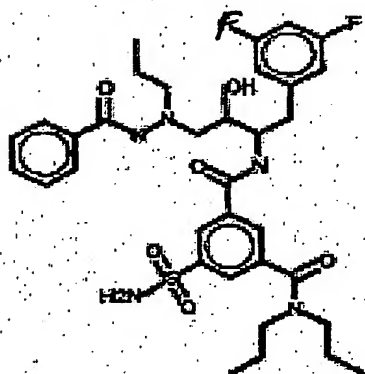
1-D12 1,3,4,4



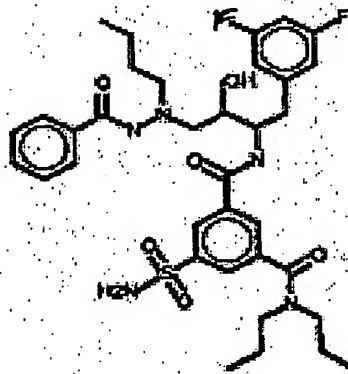
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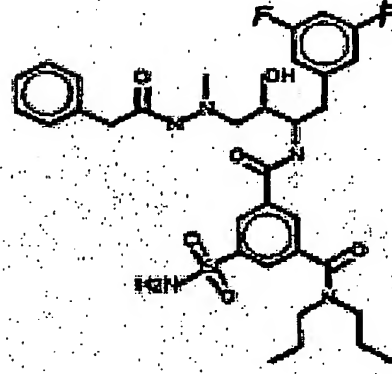
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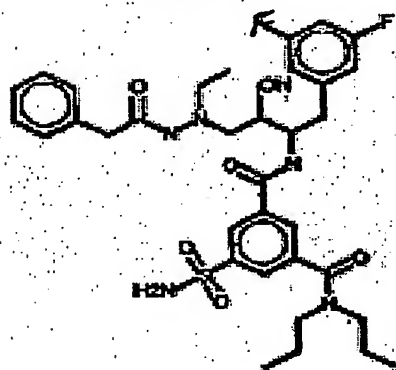
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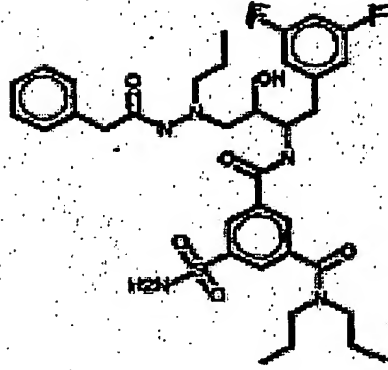
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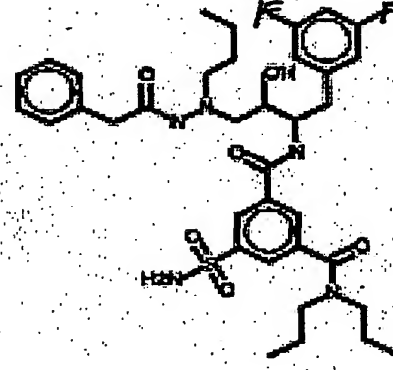
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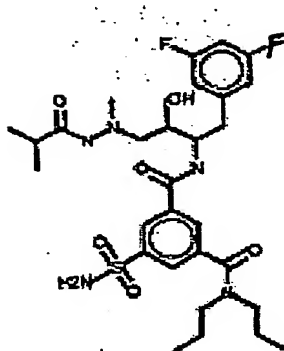
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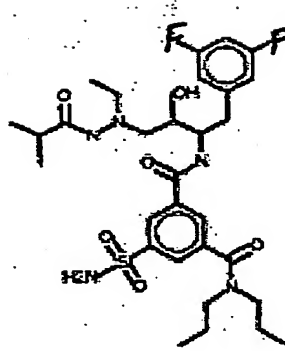
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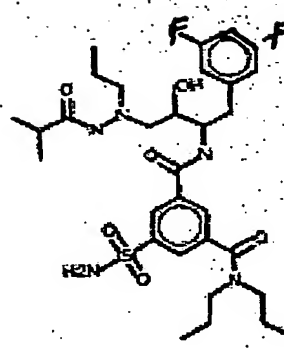
1-E8 1,4,2,4



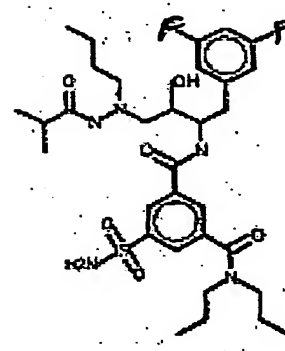
1E9 1,4,3,1



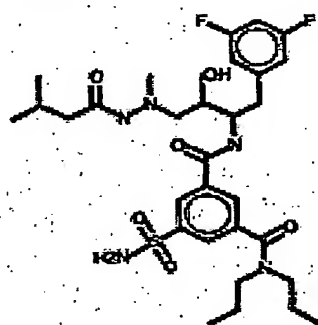
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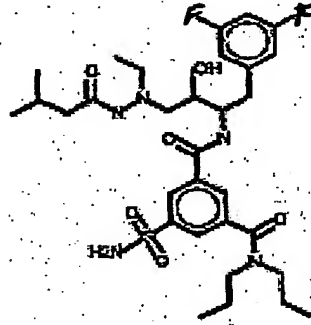
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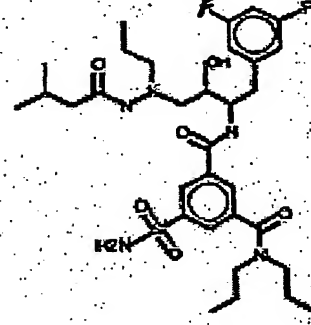
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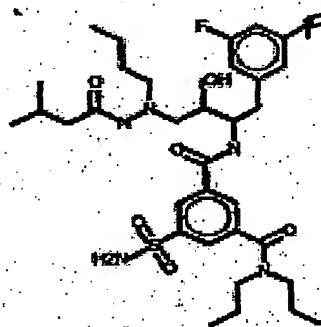
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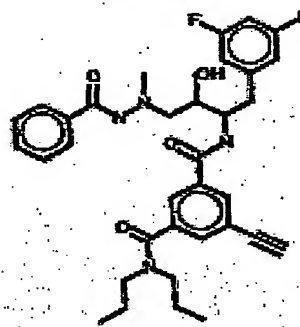
1F2 1,4,4,2



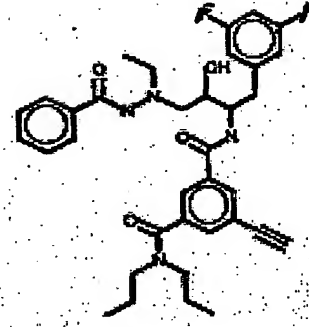
1F3 1,4,4,3



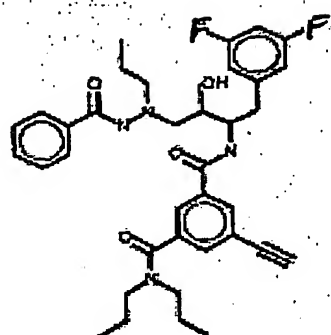
1F4 1,4,4,4



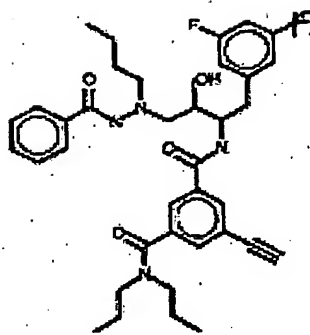
1F5 1,5,1,1



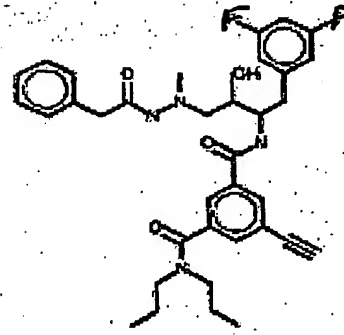
1F6 1,5,1,2



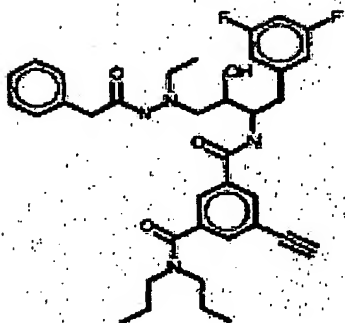
1F7 1,5,1,3



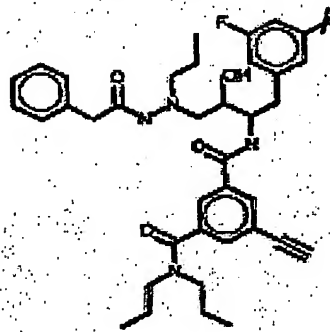
1F8 1,5,1,4



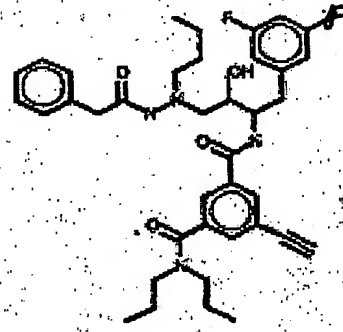
1F9 1,5,2,1



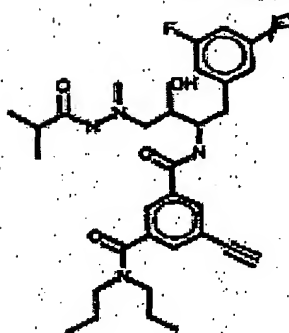
1F10 1,5,2,2



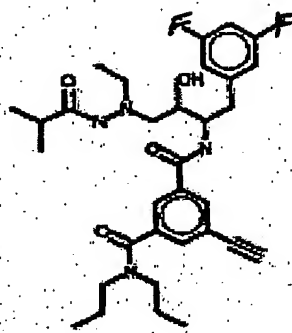
1F11 1,5,2,3



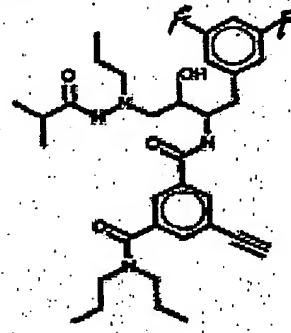
1F12 1,5,2,4



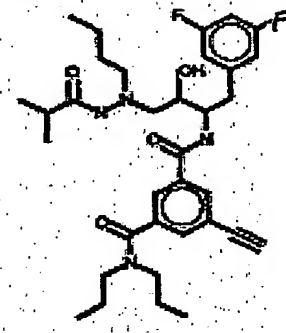
1G1 1,5,3,1



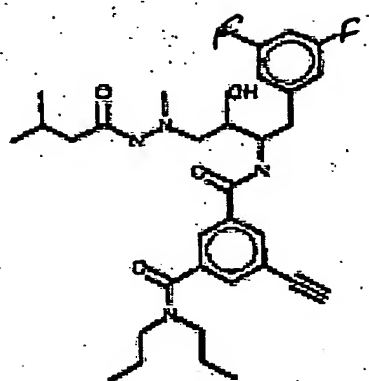
1G2 1,5,3,2



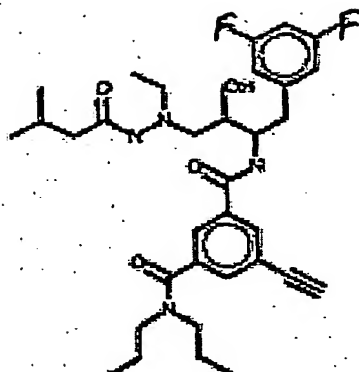
1G3 1,5,3,3



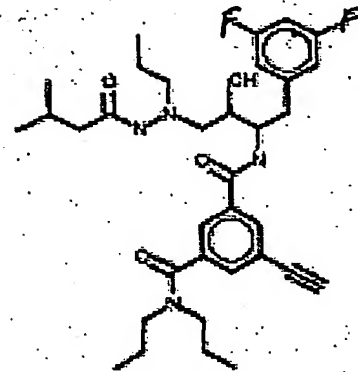
1G4 1,5,3,4



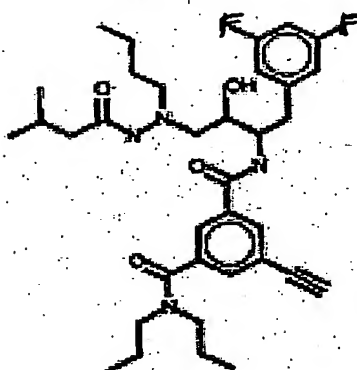
1:35 1,5,4,1



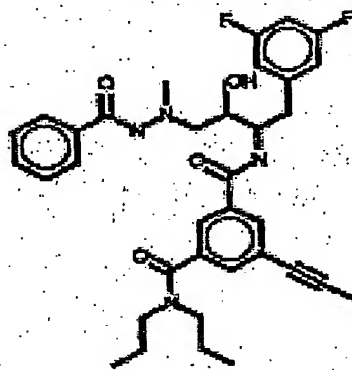
1:36 1,5,4,2



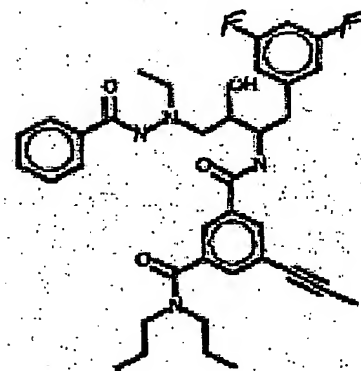
1:37 1,5,4,3



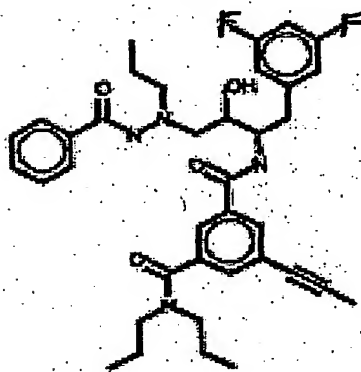
1:38 1,5,4,4



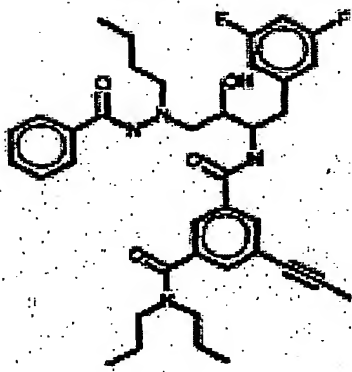
1:39 1,5,1,1



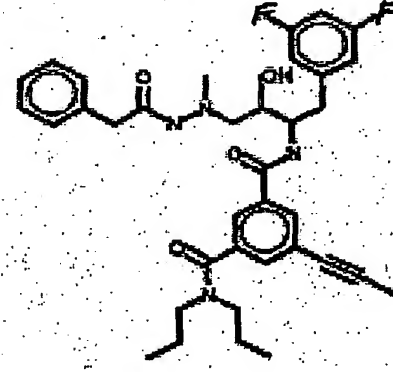
1:40 1,5,1,2



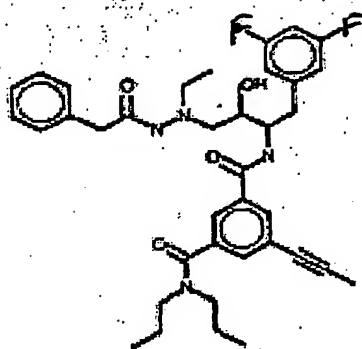
1:41 1,5,1,3



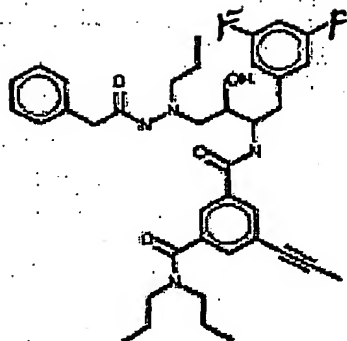
1:42 1,5,1,4



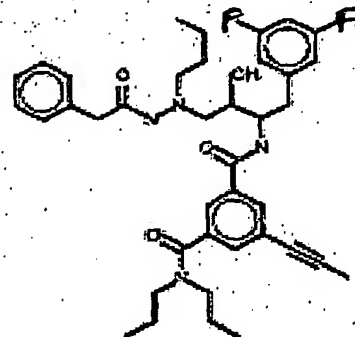
1:43 1,5,2,1



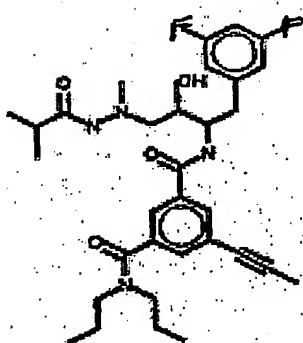
1442 1,6,2,2



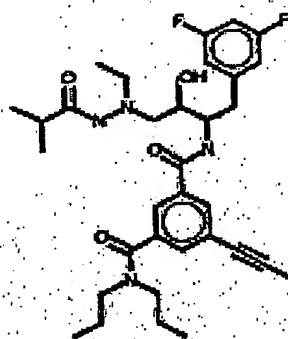
1443 1,6,2,2



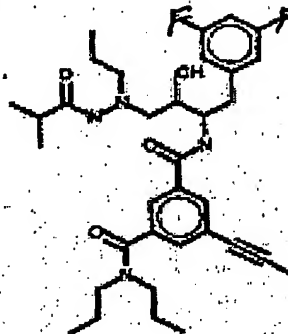
1444 1,6,2,4



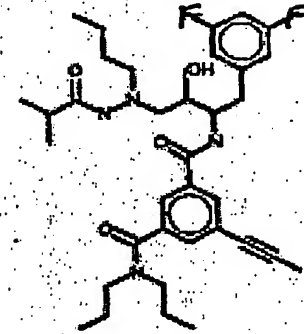
1445 1,6,3,1



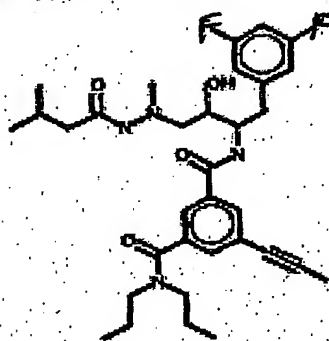
1446 1,6,3,2



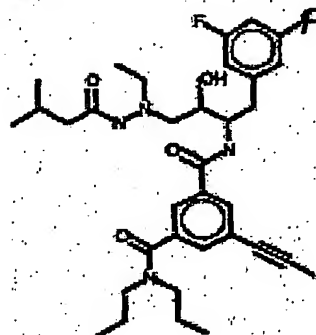
1447 1,6,3,3



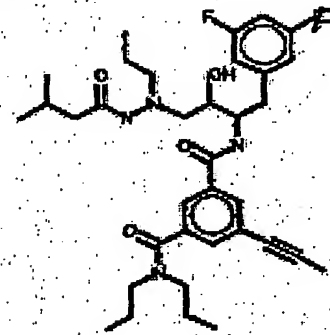
1448 1,6,3,4



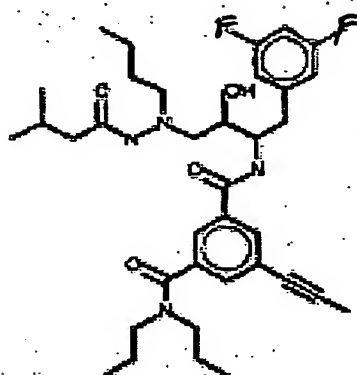
1449 1,6,4,1



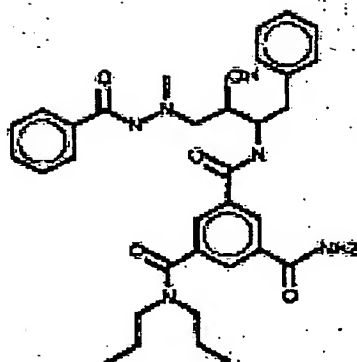
1450 1,6,4,2



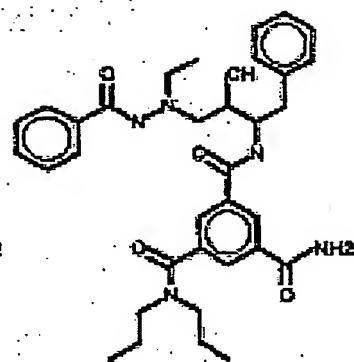
1451 1,6,4,3



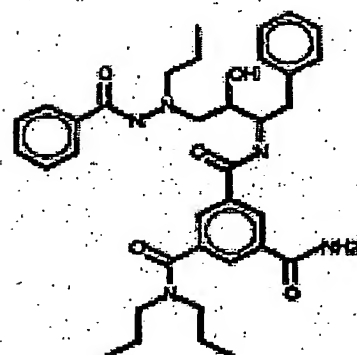
13EE2 1.6.4.4



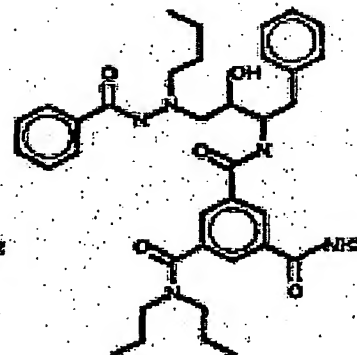
2A1 2.1.1.1



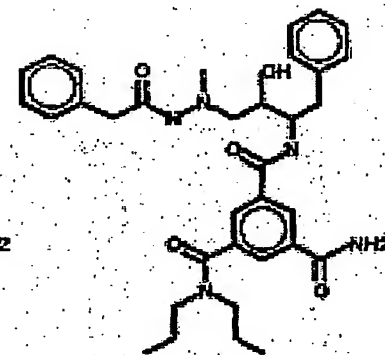
2A2 2.1.1.2



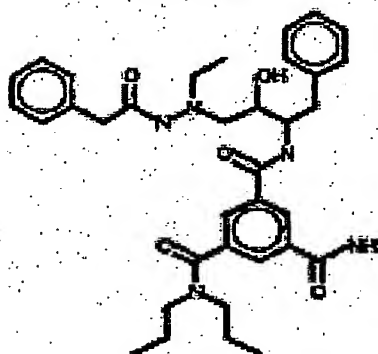
2A3 2.1.1.3



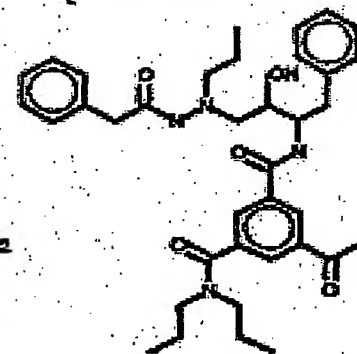
2A4 2.1.1.4



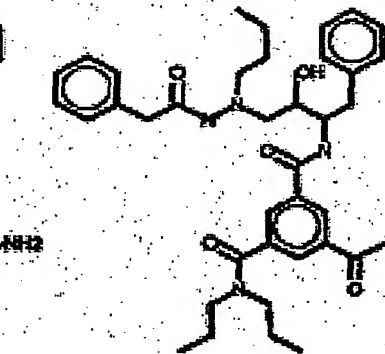
2A5 2.1.2.1



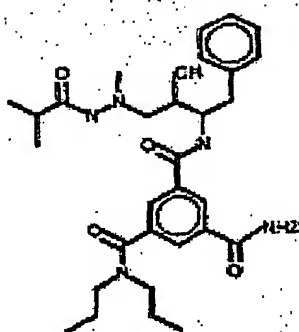
2A6 2.1.2.2



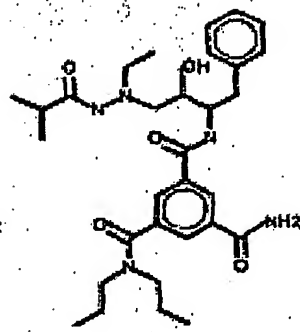
2A7 2.1.2.3



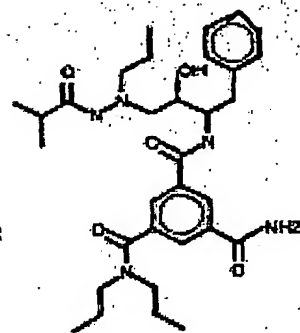
2A8 2.1.2.4



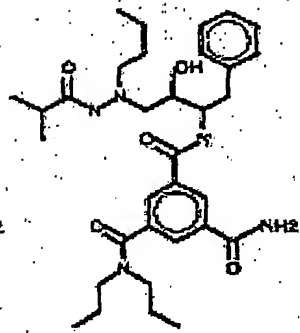
2A9 2,1,3,1



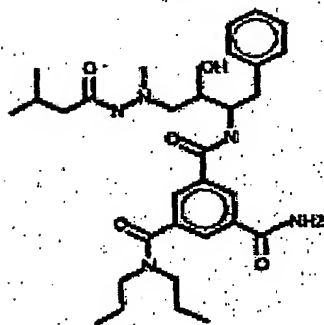
2A10 2,1,3,2



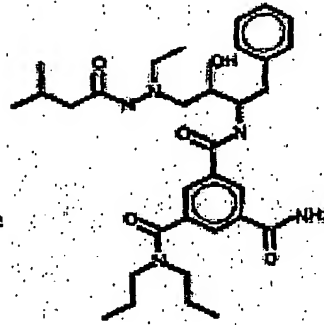
2A11 2,1,3,3



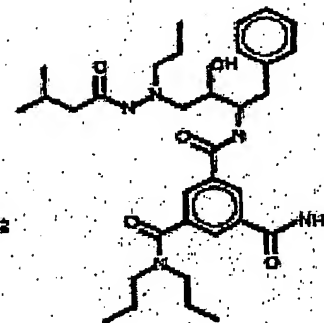
2A12 2,1,3,4



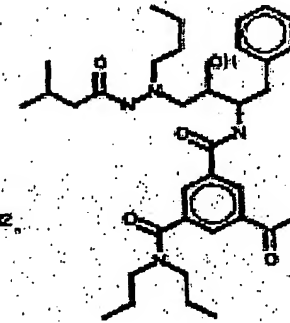
2B1 2,1,4,1



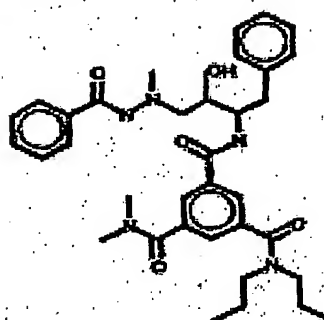
2B2 2,1,4,2



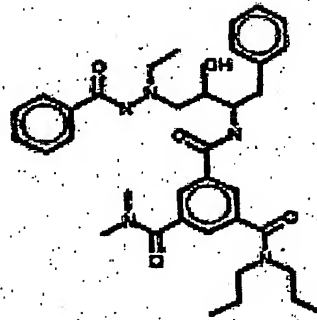
2B3 2,1,4,3



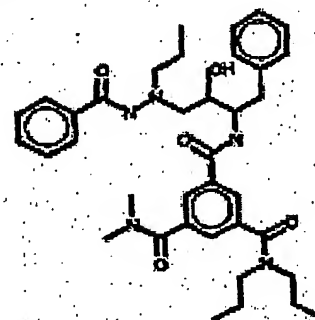
2B4 2,1,4,4



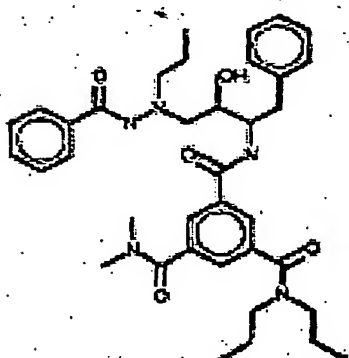
2B5 2,2,1,1



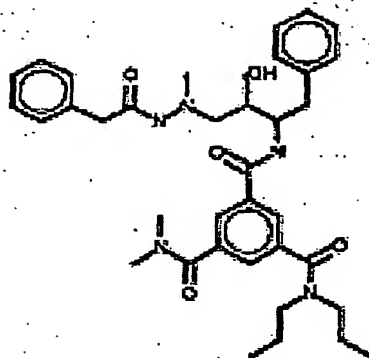
2B6 2,2,1,2



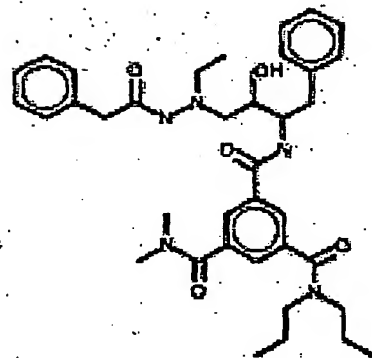
2B7 2,2,1,3



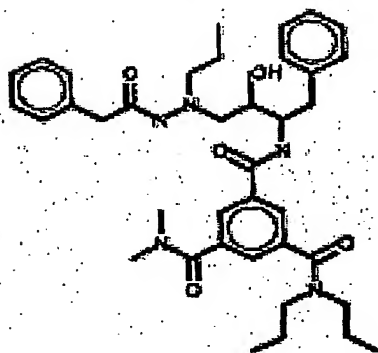
2B8 2,2,1,4



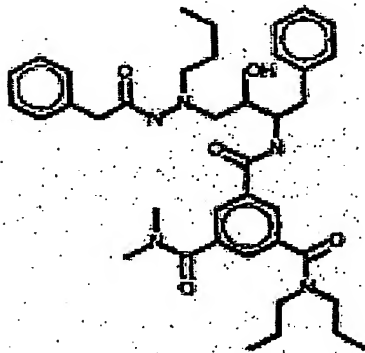
2B9 2,2,2,1



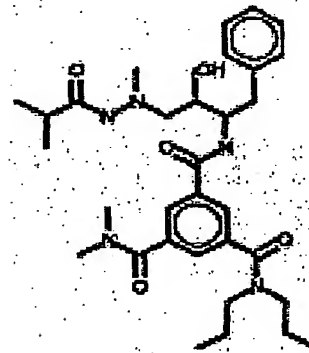
2B10 2,2,2,2



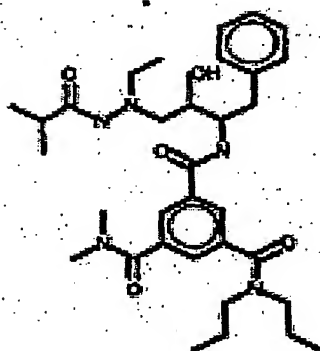
2B11 2,2,2,3



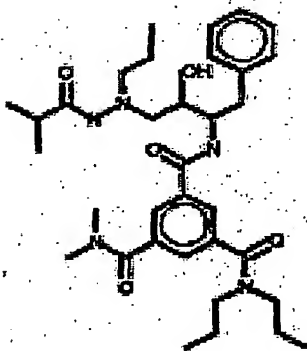
2B12 2,2,2,4



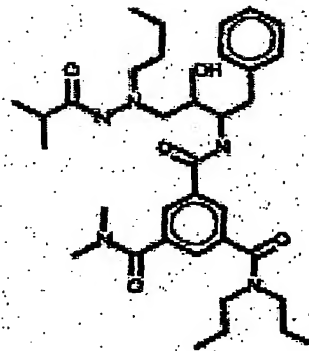
2C1 2,2,3,1



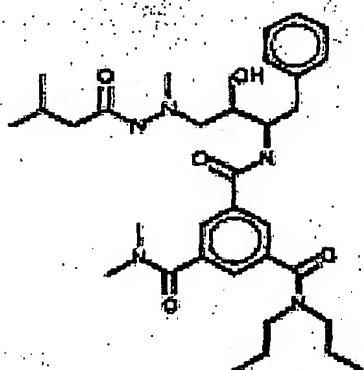
2C2 2,2,3,2



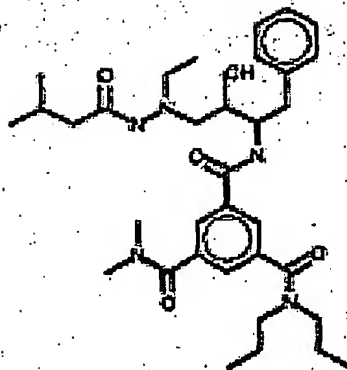
2C3 2,2,3,3



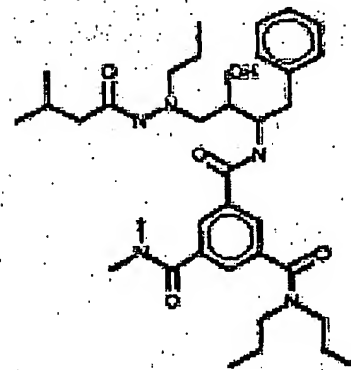
2C4 2,2,3,4



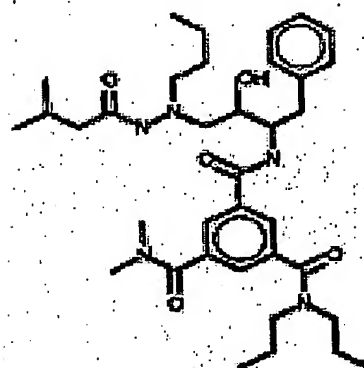
2C6 2.2.4.1



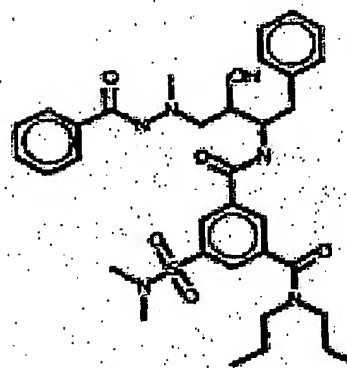
2C8 2.2.4.2



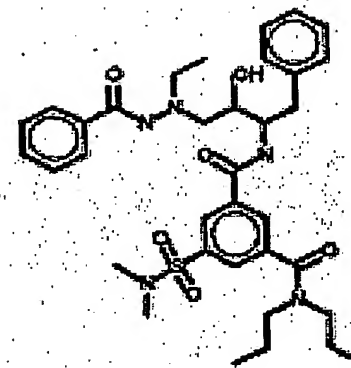
2C7 2.2.4.3



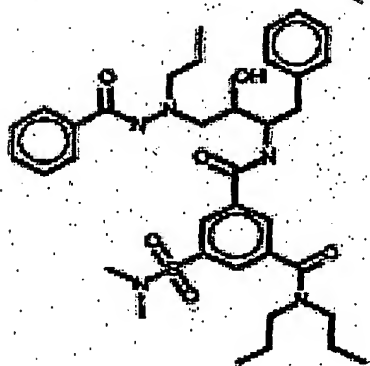
2C9 2.2.4.4



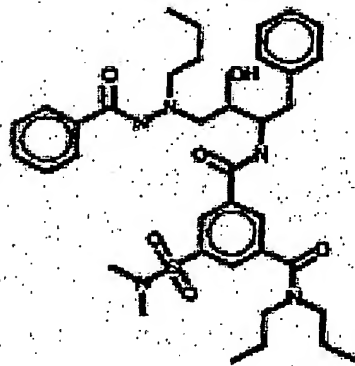
2C8 2.3.1.1



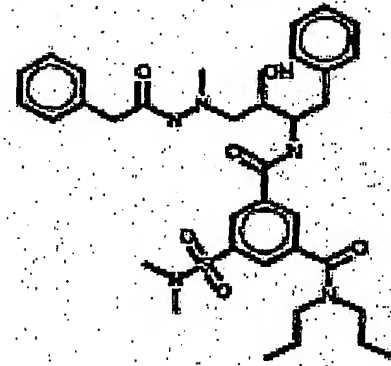
2C10 2.3.1.2



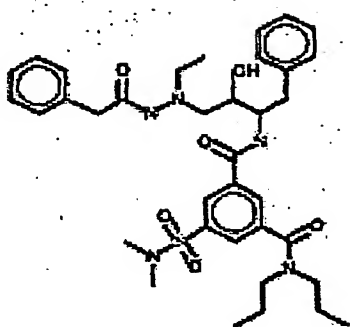
2C11 2.3.1.3



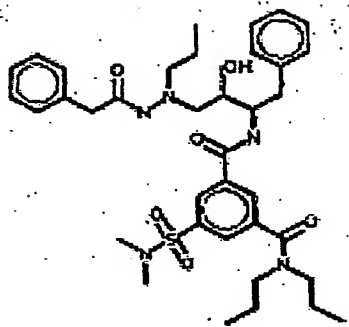
2C12 2.3.1.4



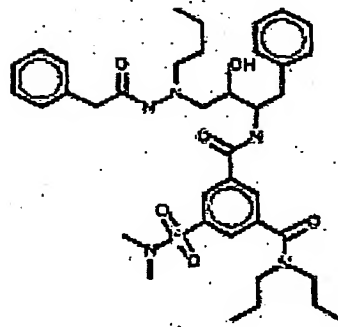
2C1 2.3.2.1



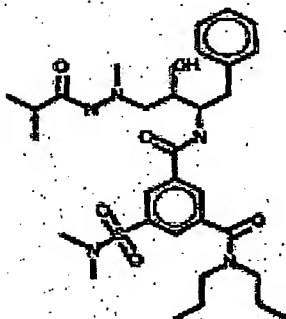
2D2 2.3.2.2



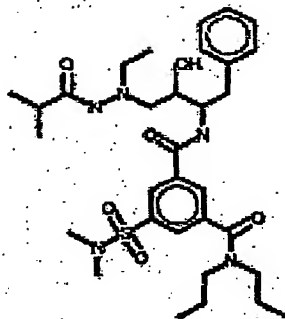
2D3 2.3.2.3



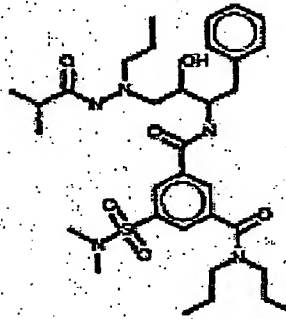
2D4 2.3.2.4



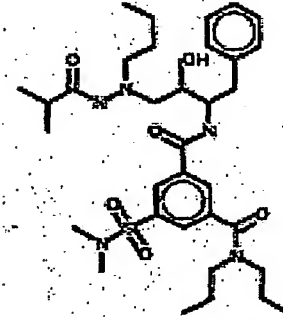
2D5 2.3.3.1



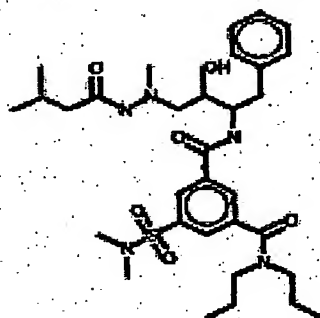
2D6 2.3.3.2



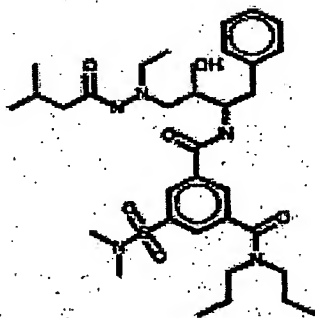
2D7 2.3.3.3



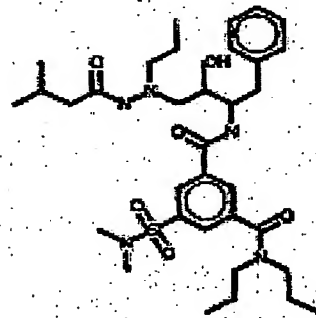
2D8 2.3.3.4



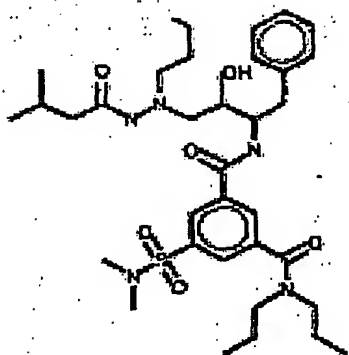
2D9 2.3.4.1



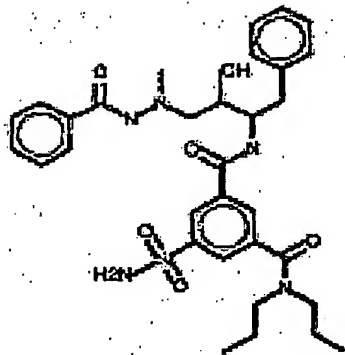
2D10 2.3.4.2



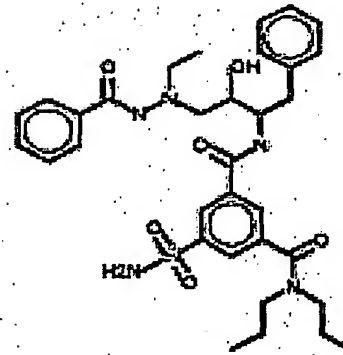
2D11 2.3.4.3



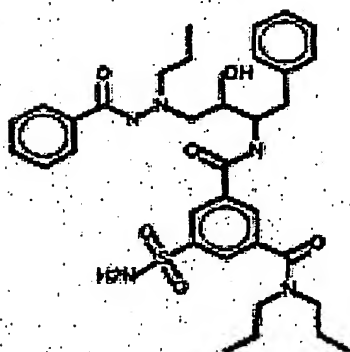
2D12 2,3,4,4



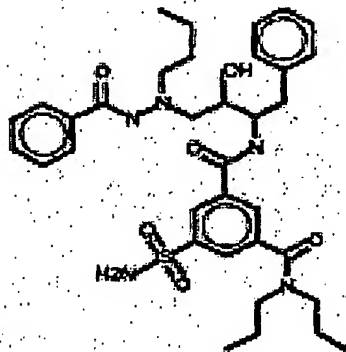
2E1 2,4,1,3



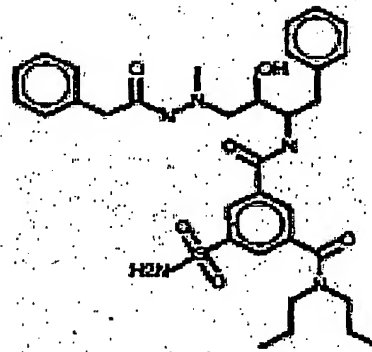
2E2 2,6,1,2



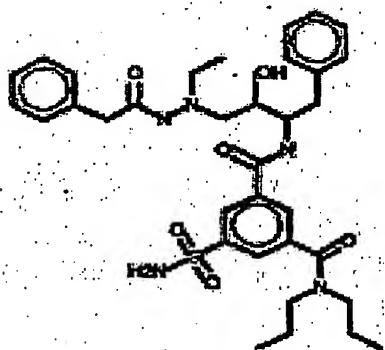
2E3 2,4,1,3



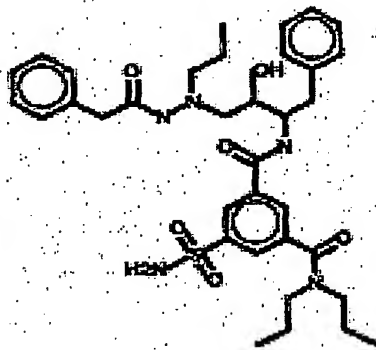
2E4 2,4,1,4



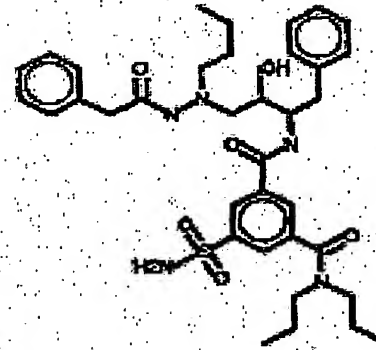
2E5 2,4,2,1



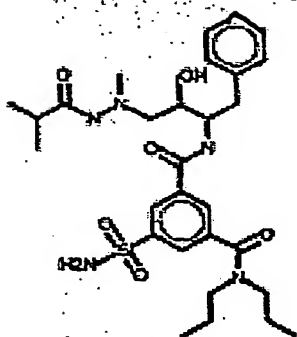
2E6 2,4,2,2



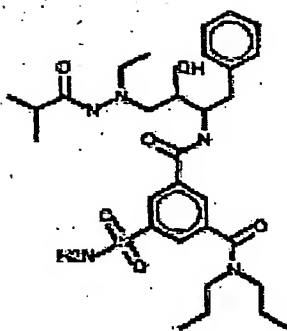
2E7 2,4,2,3



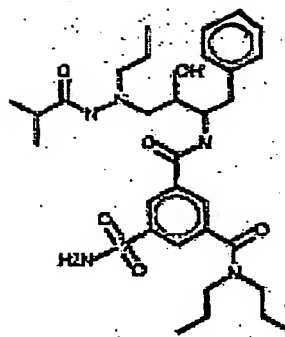
2E8 2,4,2,4



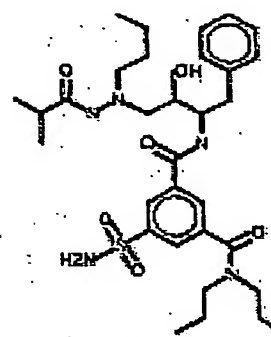
2E9 2.4.3.1



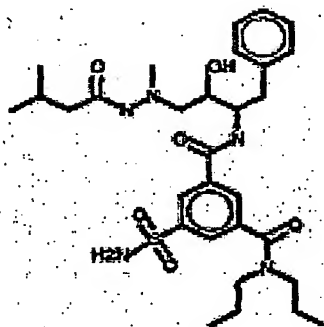
2E10 2.4.3.2



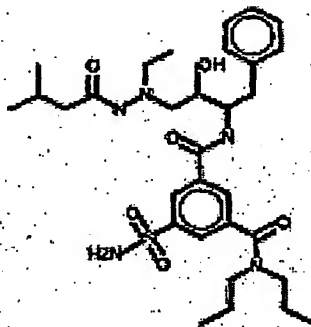
2E11 2.4.3.3



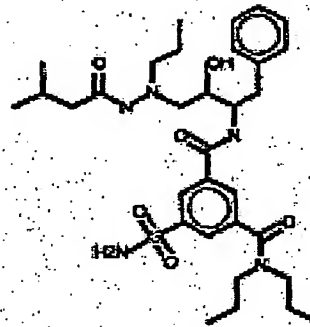
2E12 2.4.3.4



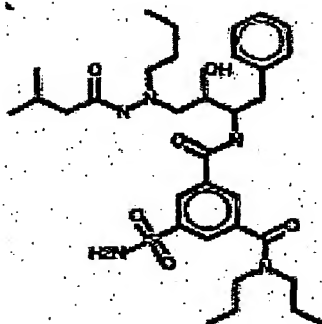
2F1 2.4.4.1



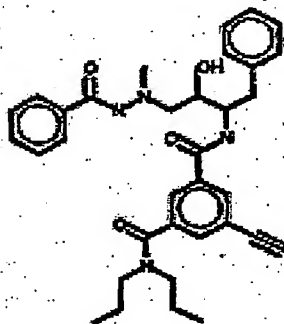
2F2 2.4.4.2



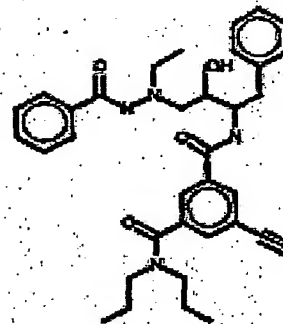
2F3 2.4.4.3



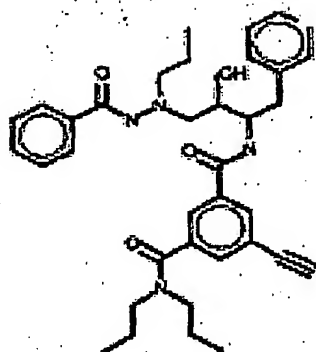
2F4 2.4.4.4



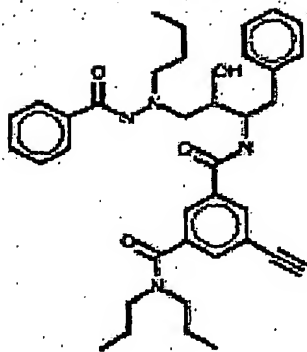
2F5 2.5.1.1



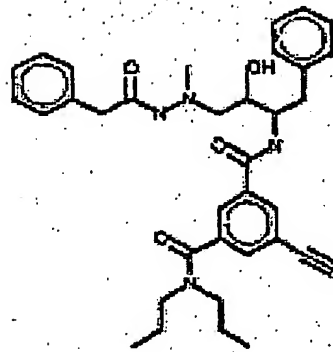
2F6 2.5.1.2



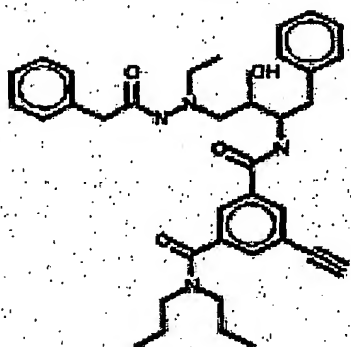
2F7 2.5.1.3



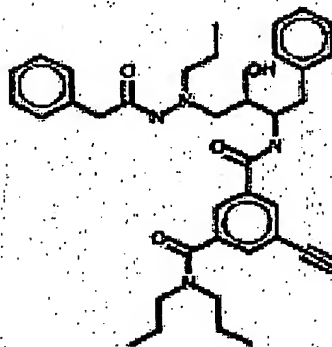
2F8 2.5.1.4



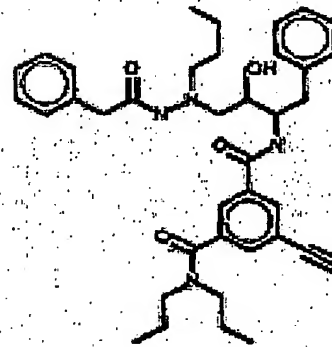
2F9 2.5.2.1



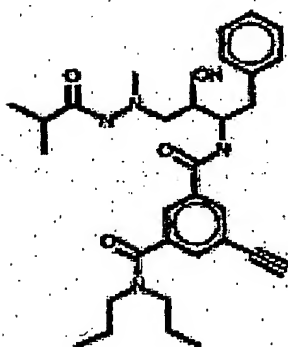
2F10 2.5.2.2



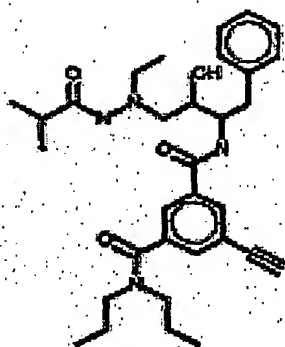
2F11 2.5.2.3



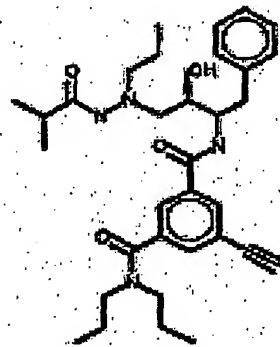
2F12 2.5.2.4



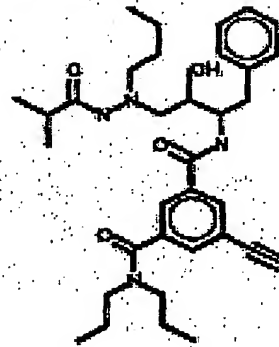
2G1 2.5.3.1



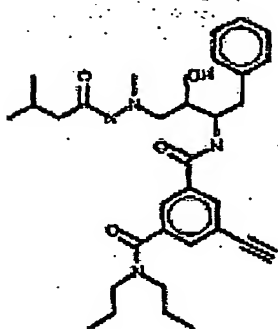
2G2 2.5.3.2



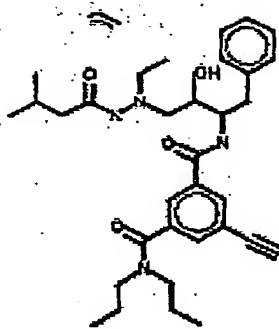
2G3 2.5.3.3



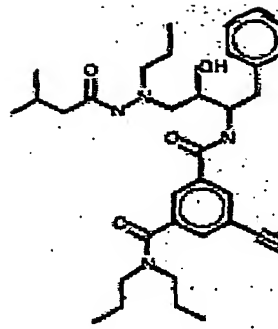
2G4 2.5.3.4



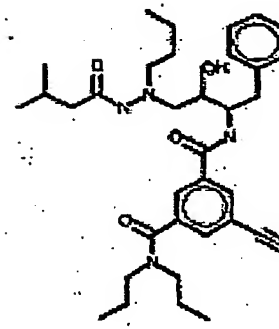
2G5 2.5.4.1



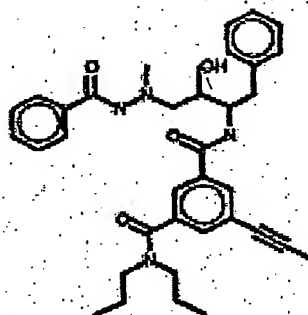
2G6 2.5.4.2



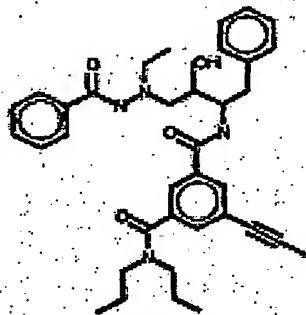
2G7 2.5.4.3



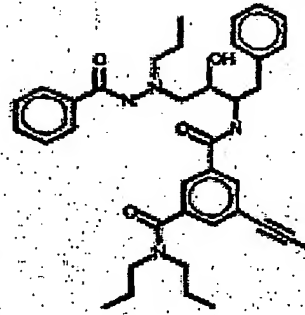
2G8 2.5.4.4



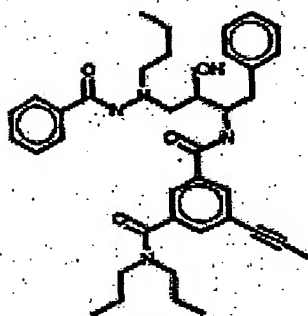
2G9 2.5.1.1



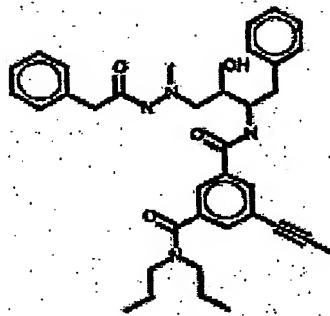
2G10 2.5.1.2



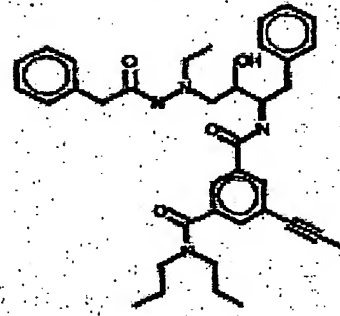
2G11 2.5.1.3



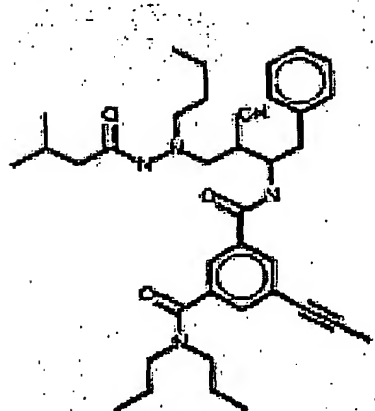
2G12 2.5.1.4



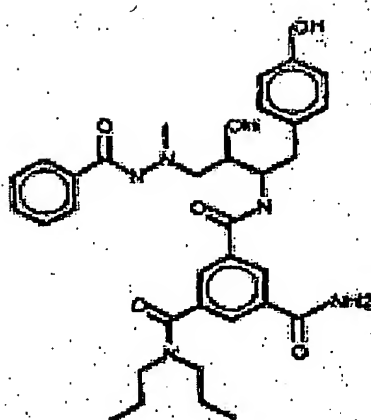
2H1 2.5.2.1



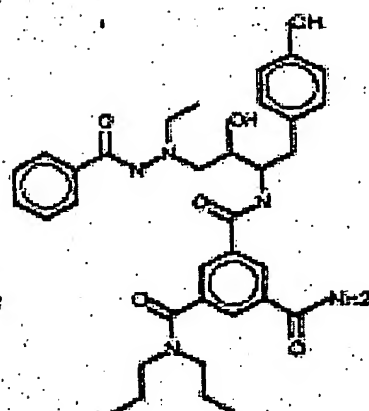
2H2 2.5.2.2



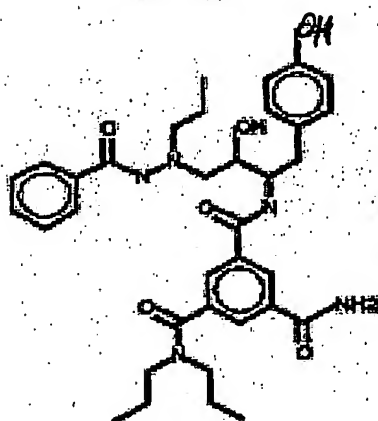
2H12 2,0,4,4



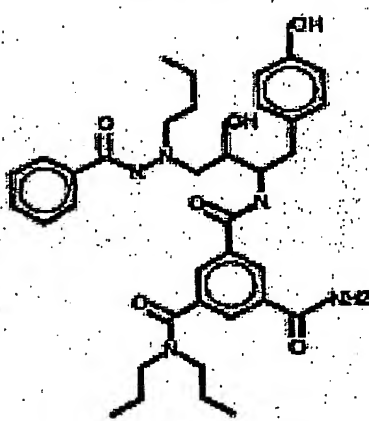
3A1 2,1,2,1



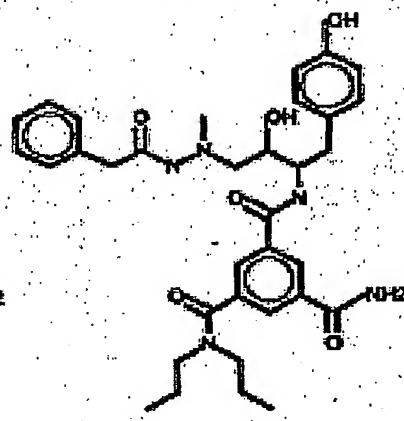
3A2 3,1,1,2



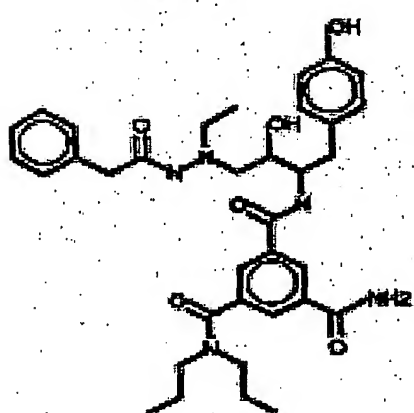
3A3 0,1,1,3



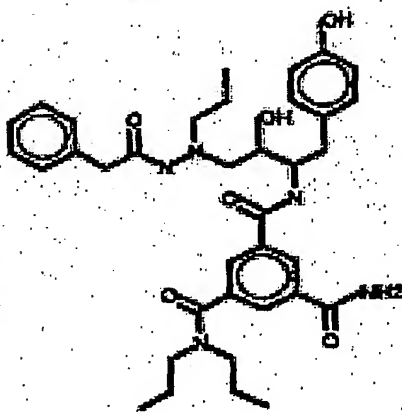
3A4 3,1,1,4



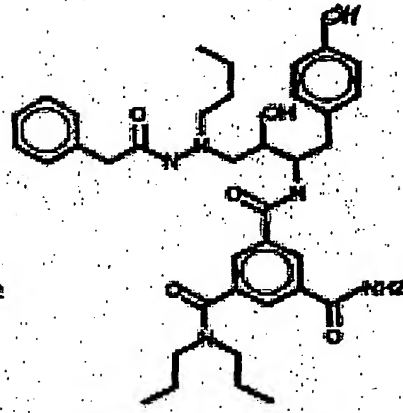
3A5 3,1,2,1



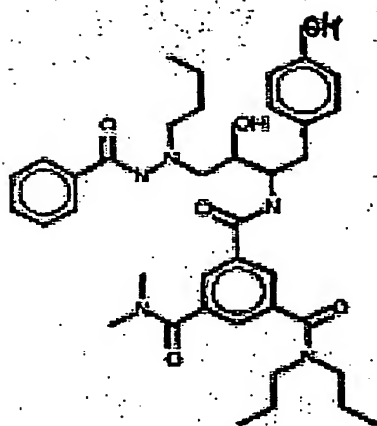
3A6 3,1,2,2



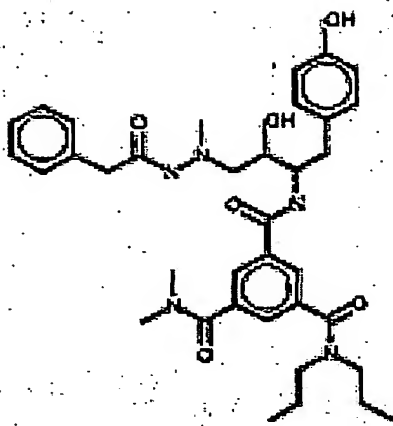
3A7 3,1,2,3



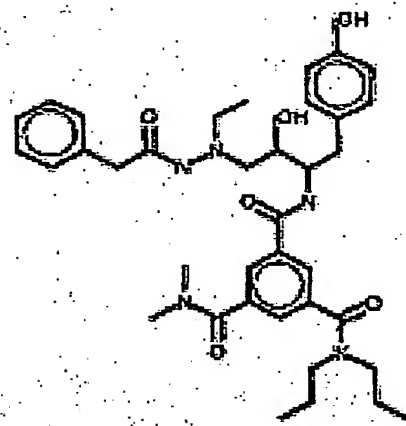
3A8 3,1,2,4



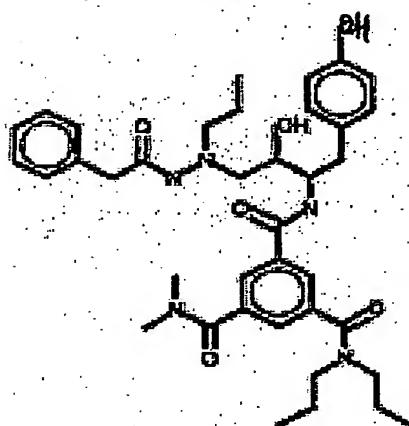
3.88 3.2.1.4



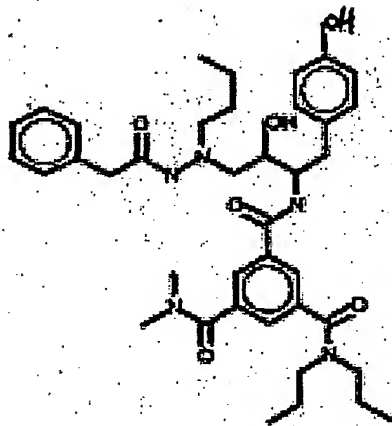
3.89 3.2.2.1



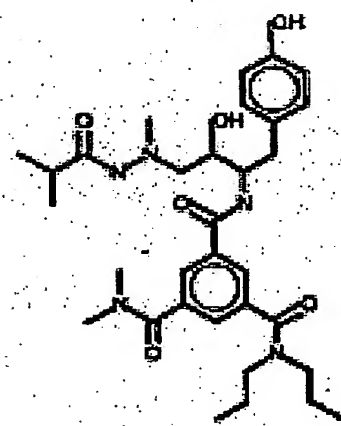
3.89 3.2.2.2



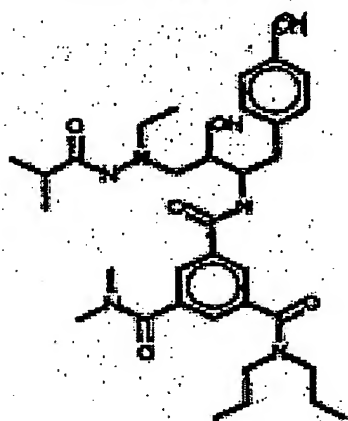
3.811 3.2.2.3



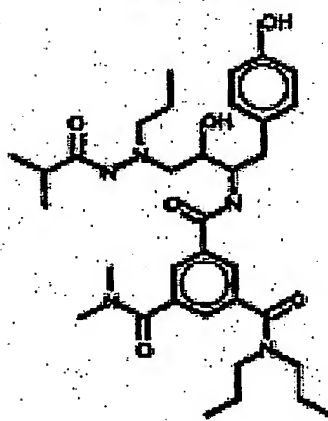
3.812 3.2.2.4



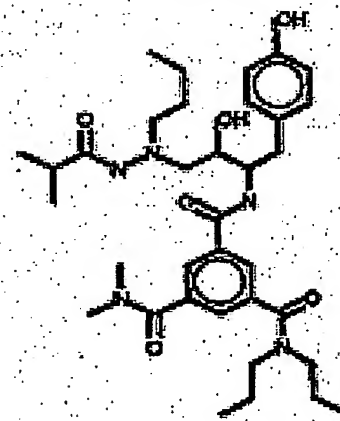
3.81 3.2.3.1



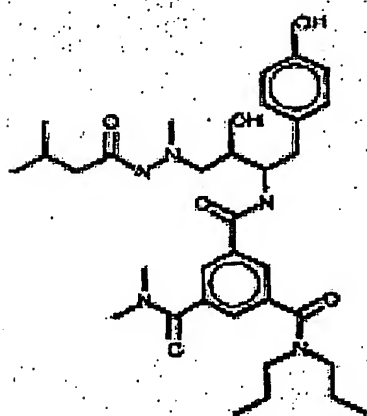
3.82 3.2.3.2



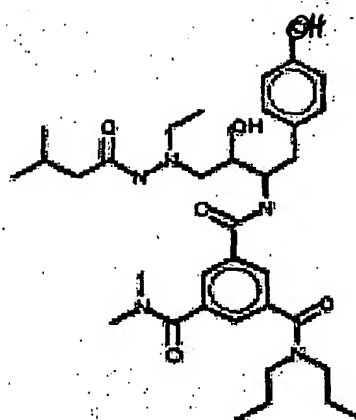
3.83 3.2.3.3



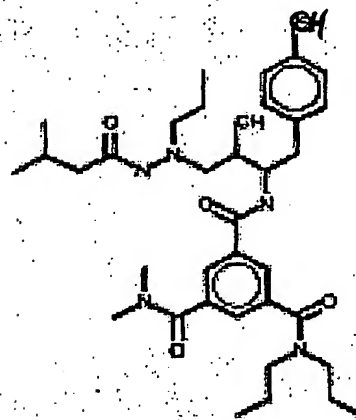
3.84 3.2.3.4



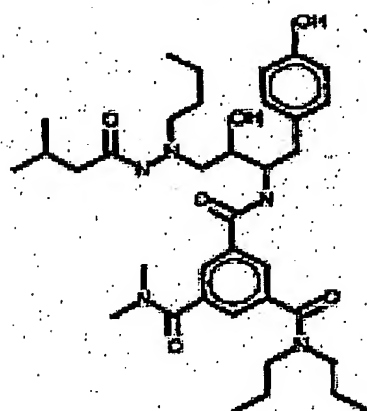
3C6 3.2.4.1



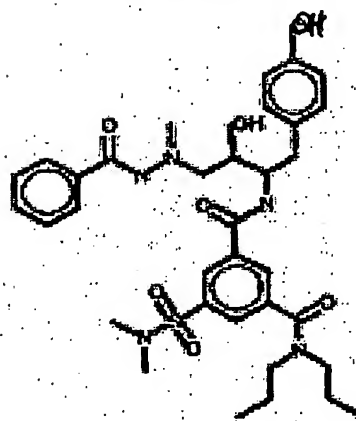
3C8 3.2.4.2



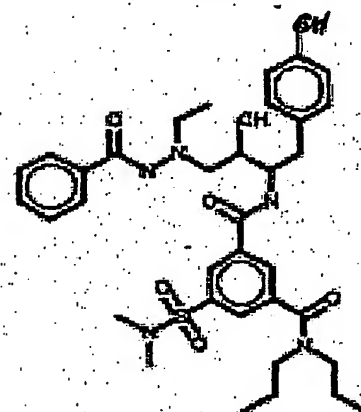
3C7 3.2.4.3



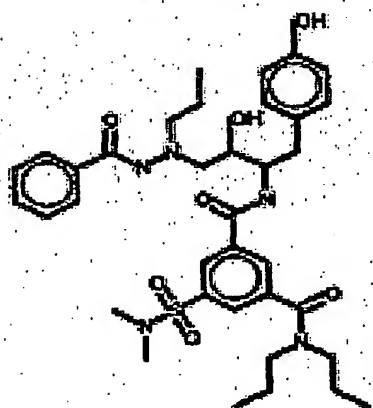
3C9 3.2.4.4



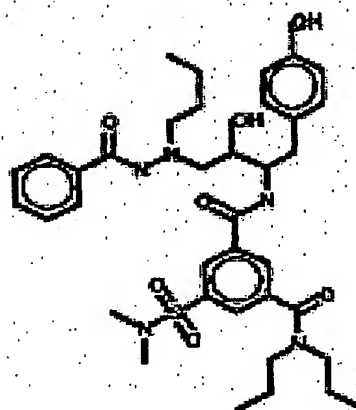
3C9 3.2.1.1



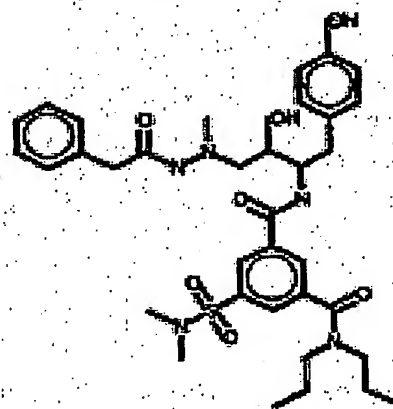
3C10 3.2.1.2



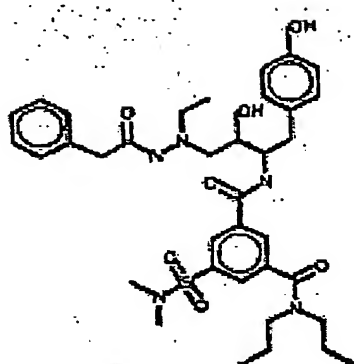
3C11 3.2.1.3



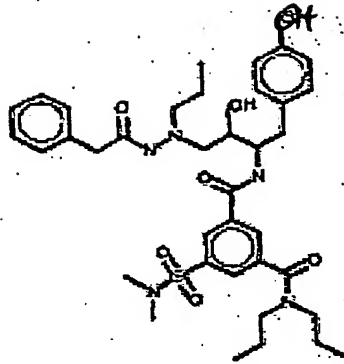
3C12 3.2.1.4



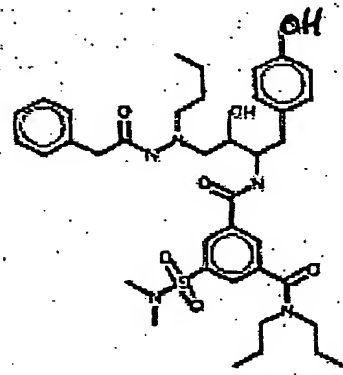
3D1 3.2.2.1



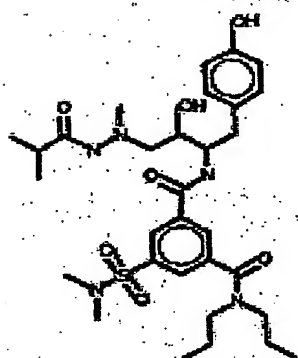
3D2 3.3.2.2



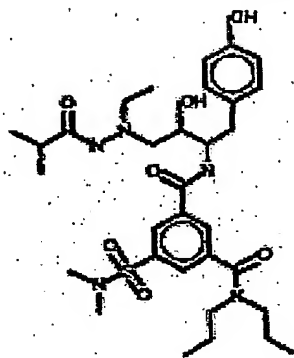
3D3 3.3.2.3



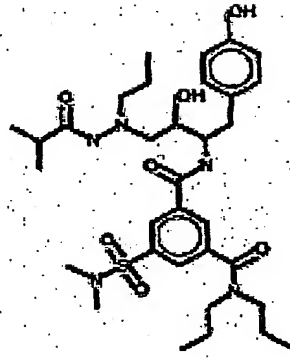
3D4 3.3.2.4



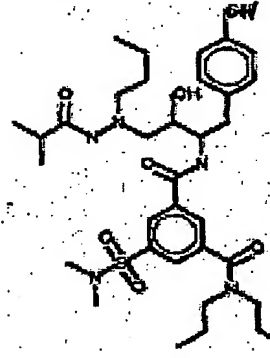
3D5 3.3.3.1



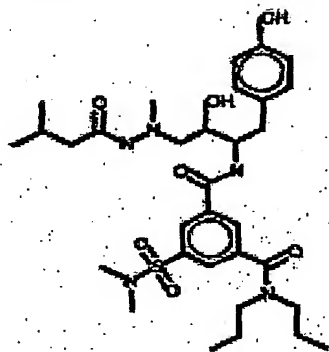
3D6 3.3.3.2



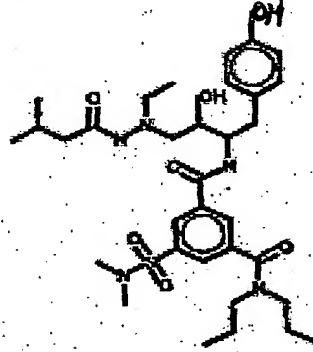
3D7 3.3.3.3



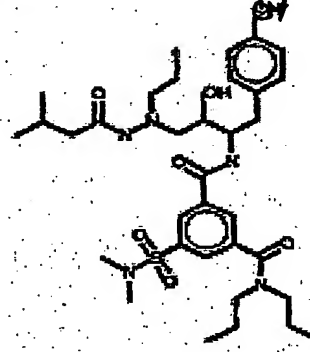
3D8 3.3.3.4



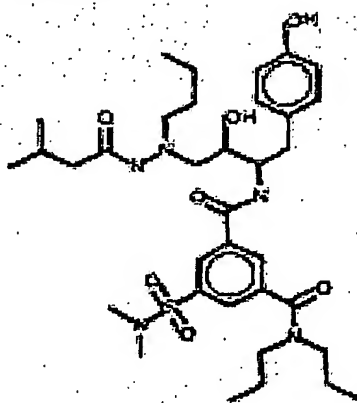
3D9 3.3.4.1



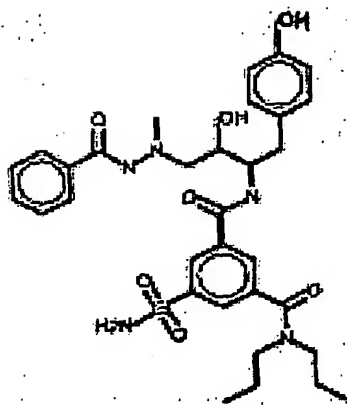
3D10 3.3.4.2



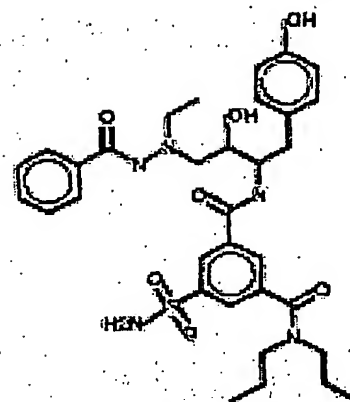
3D11 3.3.4.3



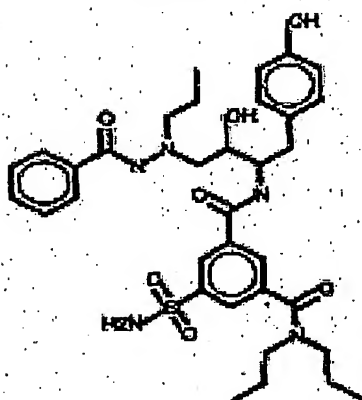
3D12 3,3,4,4



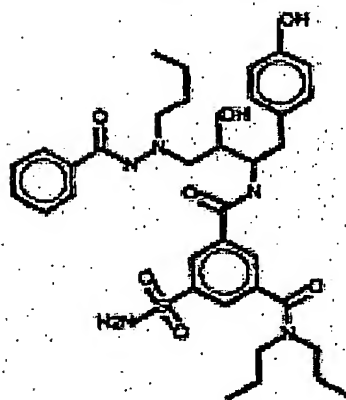
3E1 3,4,1,1



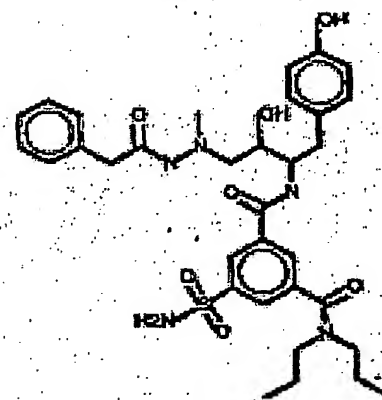
3E2 3,4,1,2



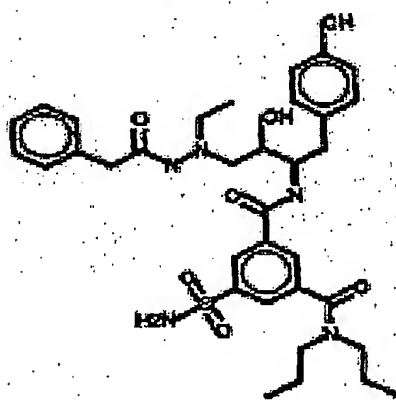
3E3 3,4,1,3



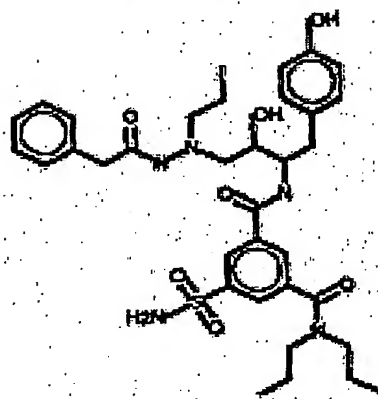
3E4 3,4,1,4



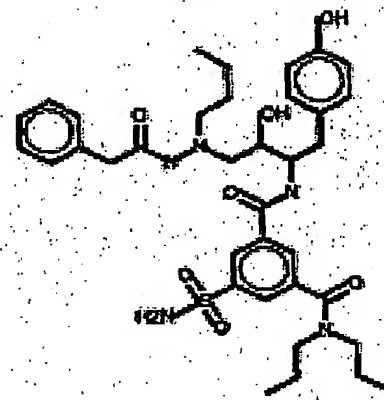
3E5 3,4,2,1



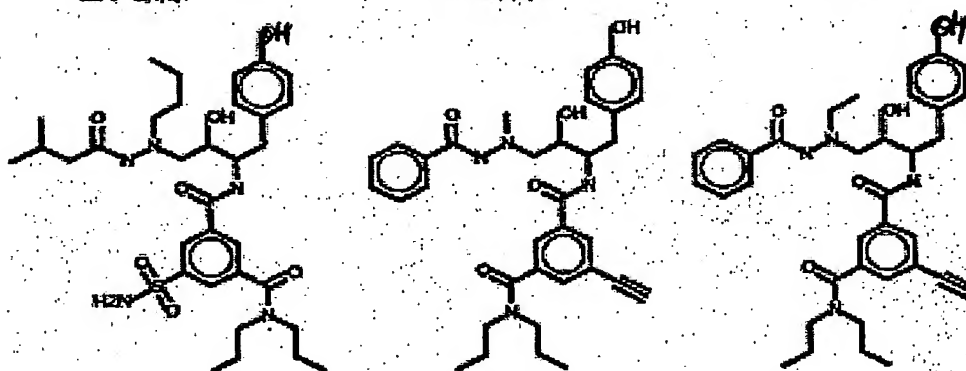
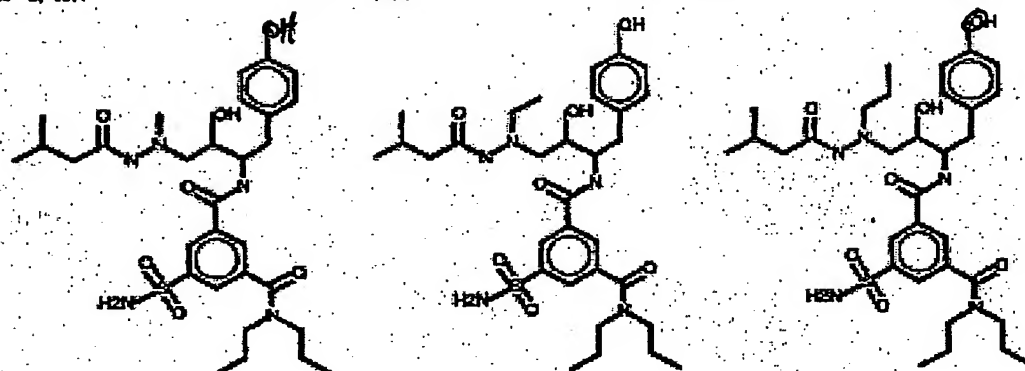
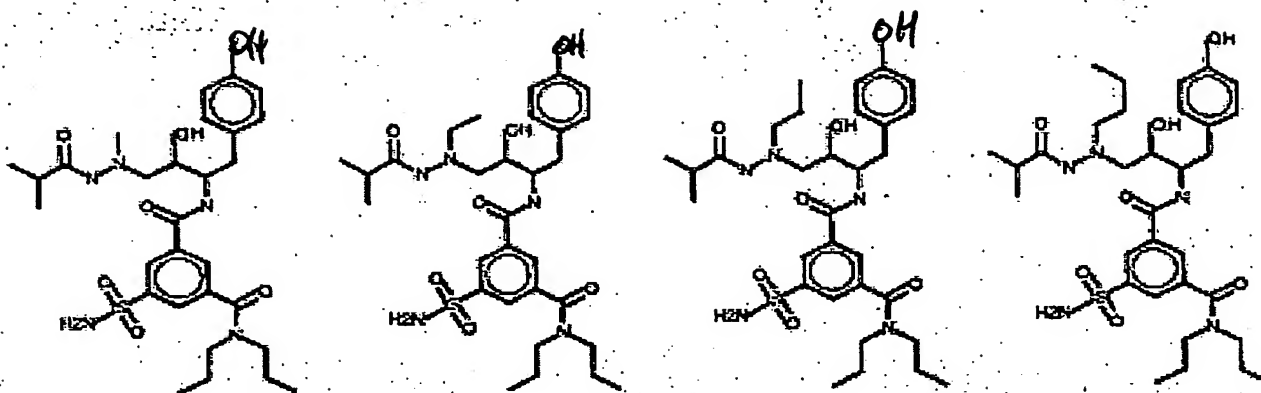
3E6 3,4,2,2

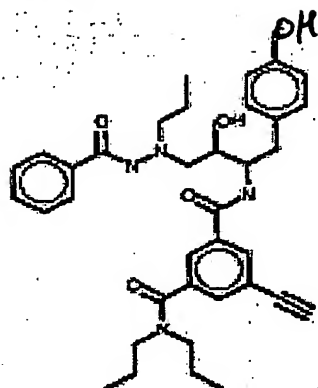


3E7 3,4,2,3

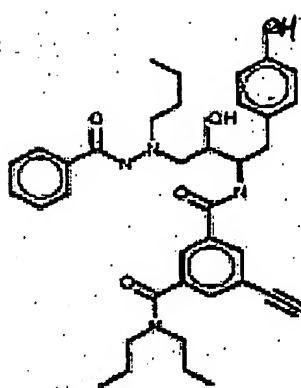


3E8 3,4,2,4

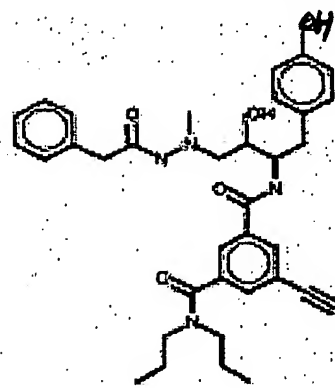




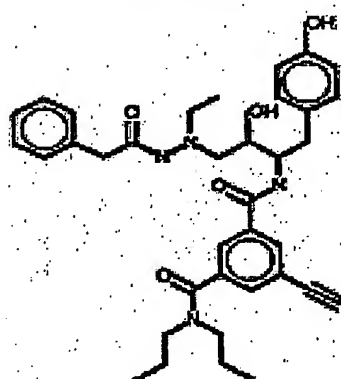
3-F7 3.5.1.3



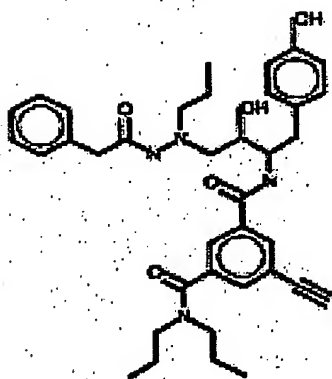
3-F8 3.5.1.4



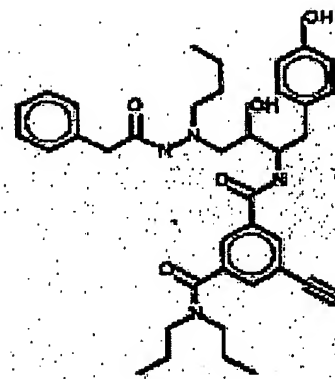
3-F9 3.5.2.1



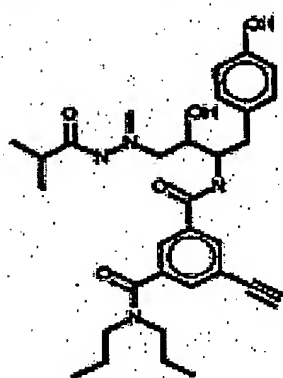
3-F10 3.5.2.2



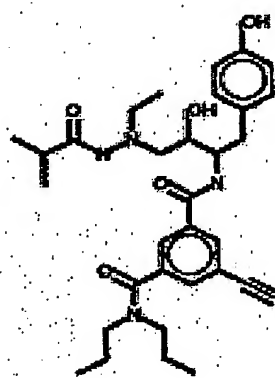
3-F11 3.5.2.3



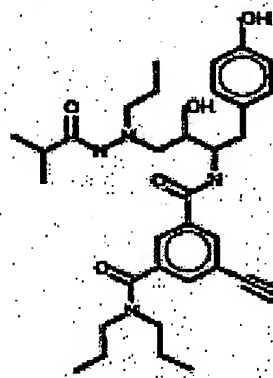
3-F12 3.5.2.4



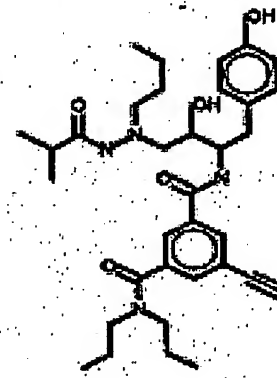
3-G1 3.5.3.1



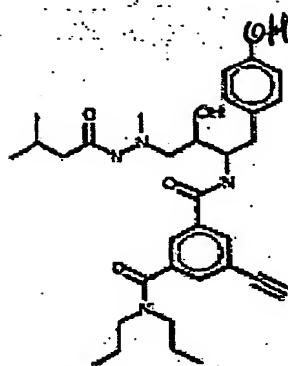
3-G2 3.5.3.2



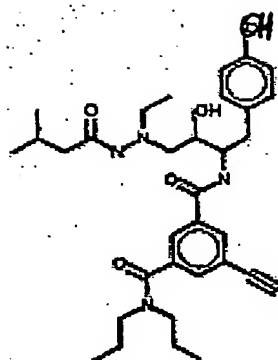
3-G3 3.5.3.3



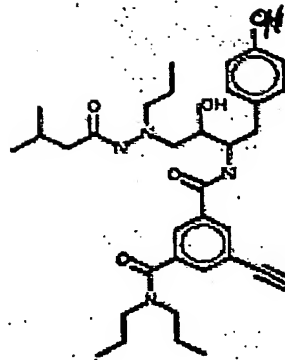
3-G4 3.5.3.4



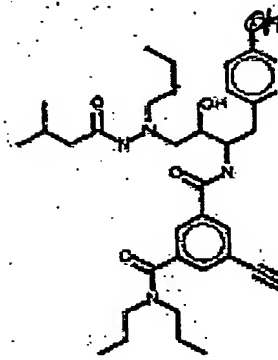
3G5 3.5.4.1



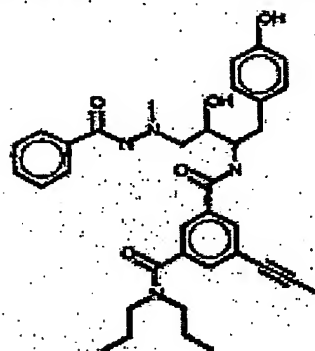
3G6 3.5.4.2



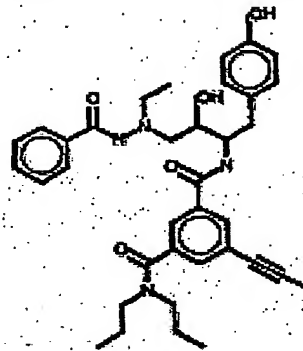
3G7 3.5.4.3



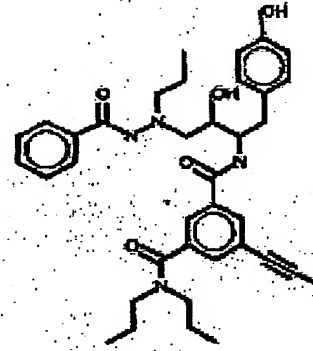
3G8 3.5.4.4



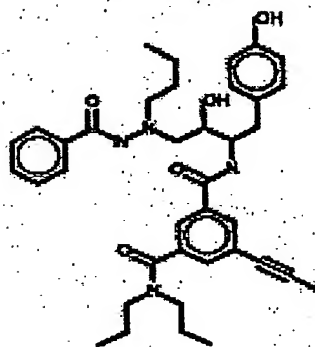
3G9 3.5.1.1



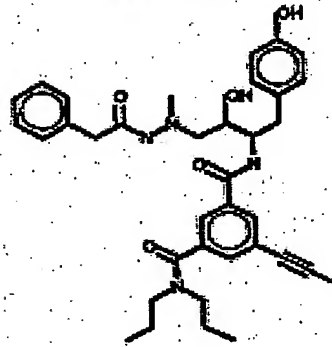
3G10 3.5.1.2



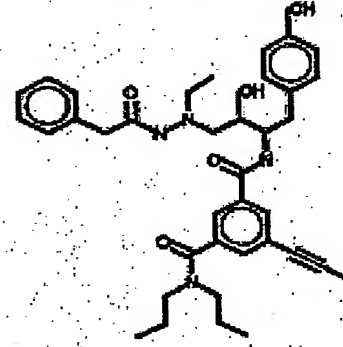
3G11 3.5.1.3



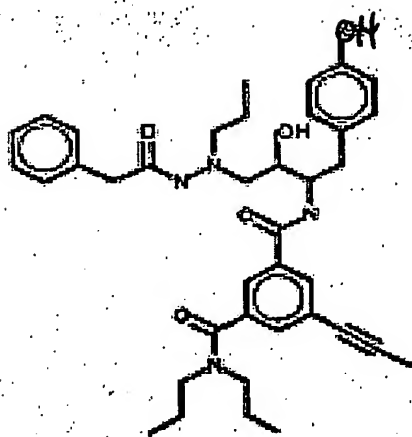
3G12 3.5.1.4



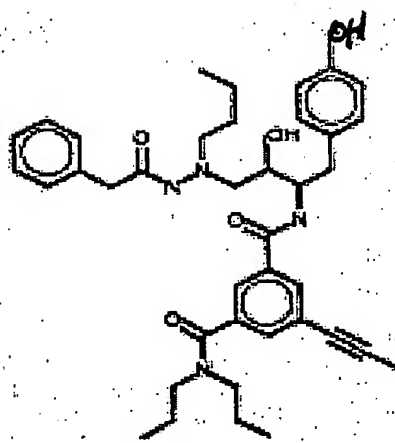
3G13 3.5.2.1



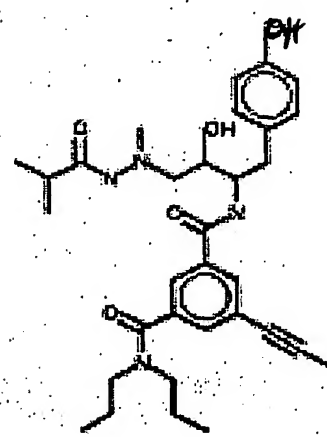
3G14 3.5.2.2



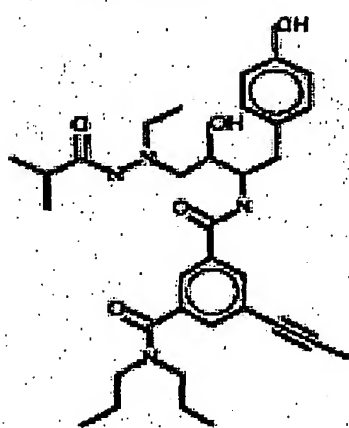
3H3 3,5,2,3



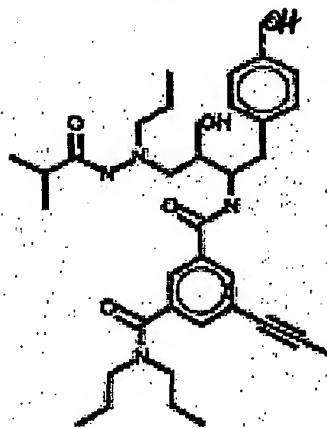
3H4 3,5,2,4



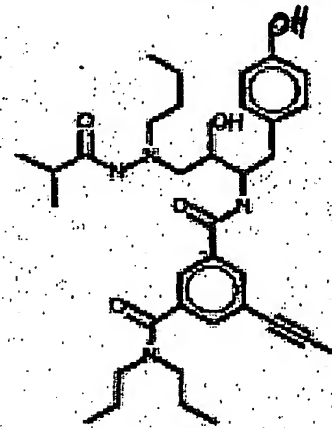
3H5 3,5,3,1



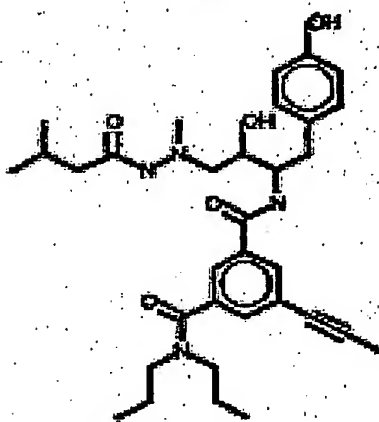
3H6 3,5,3,2



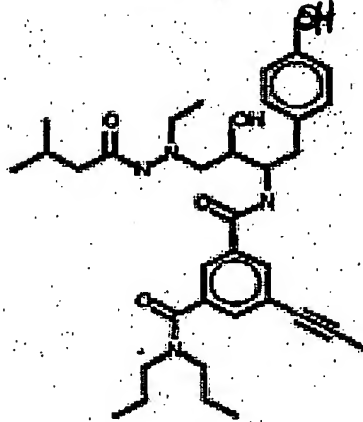
3H7 3,5,3,3



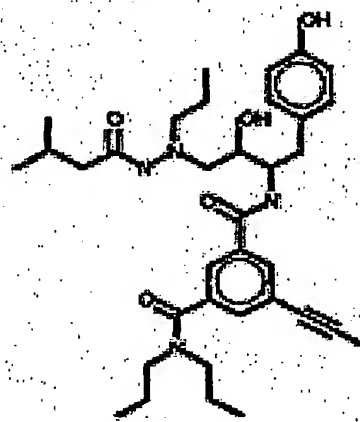
3H8 3,5,3,4



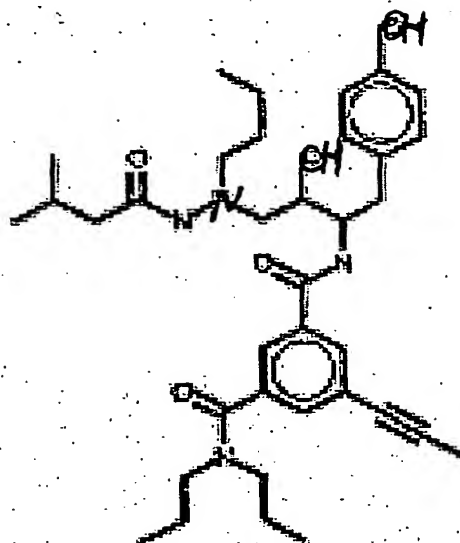
3H9 3,5,4,1



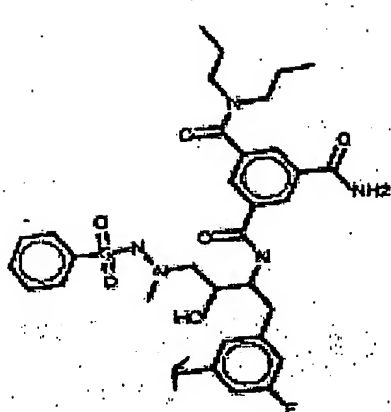
3H10 3,5,4,2



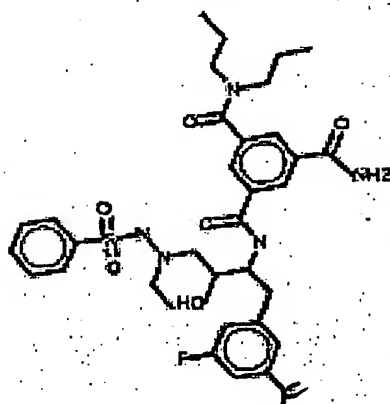
3H11 3,5,4,3



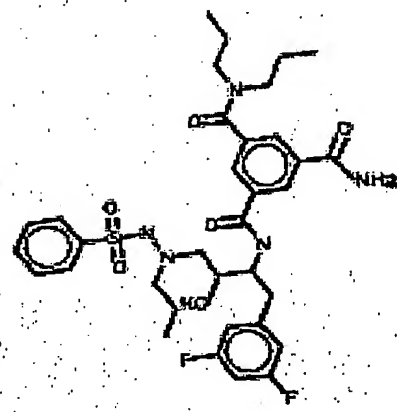
3H12 3A,4A

Chemical library "FDA581-2"
With location, precursor

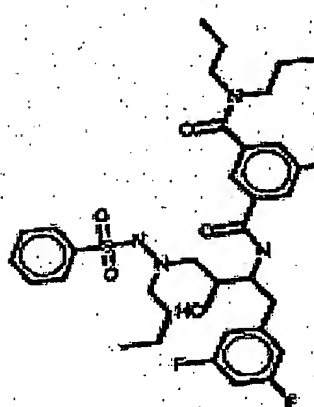
1A1 1,1,1,1



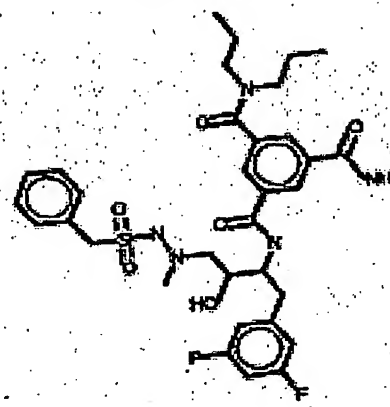
1A2 1,1,1,2



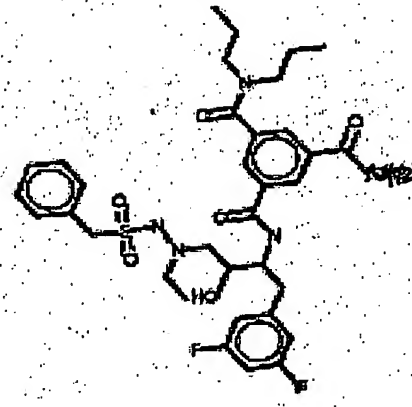
1A3 1,1,1,3



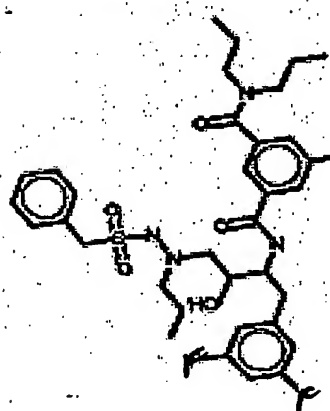
1A4 1,1,1,4



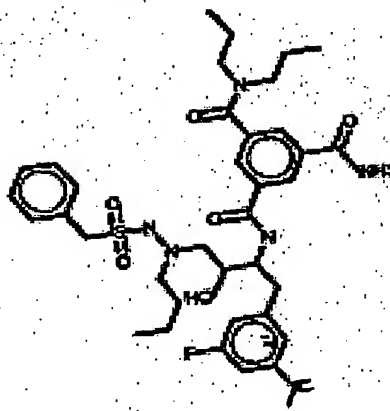
1A5 1,1,2,1



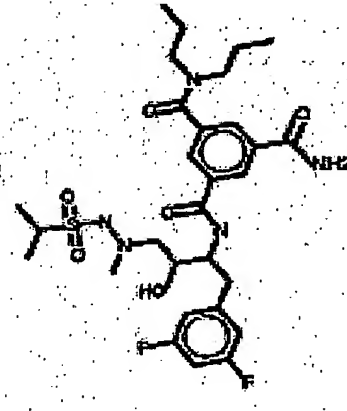
1A6 1,1,2,2



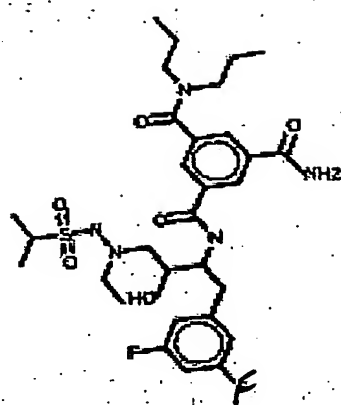
1A7 1,1,2,3



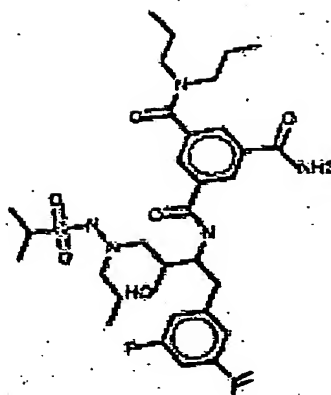
1A8 1,1,2,4



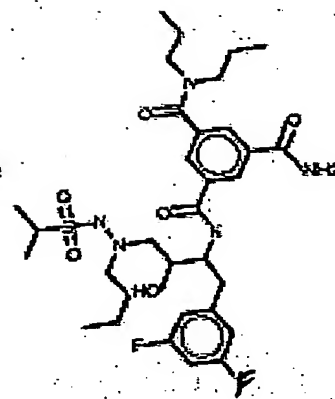
1A9 1,1,3,1



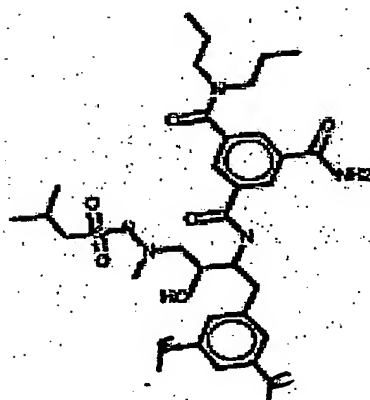
1A10 1,1,3,2



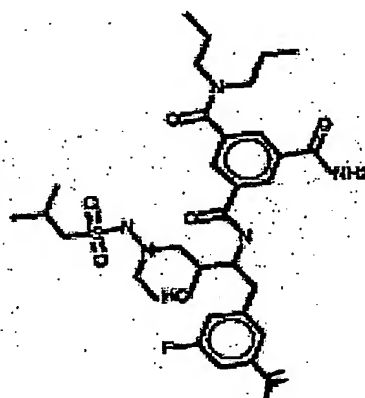
1A11 1,1,3,3



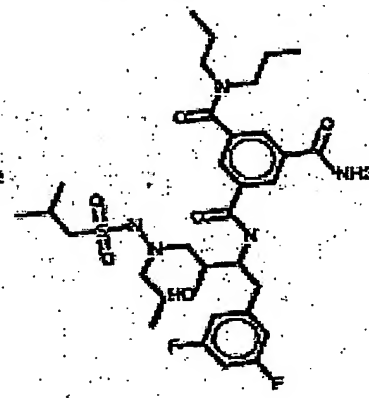
1A12 1,1,3,4



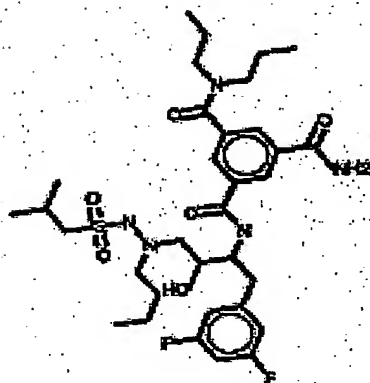
1B1 1,1,4,1



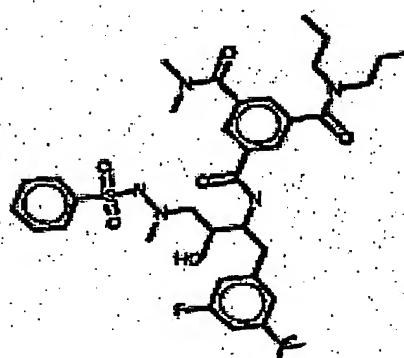
1B2 1,1,4,2



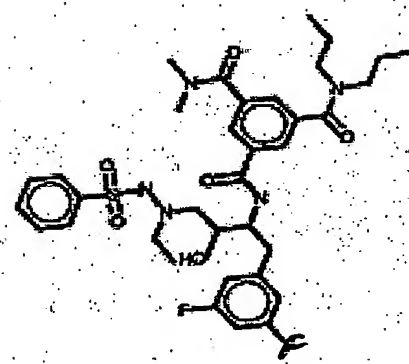
1B3 1,1,4,3



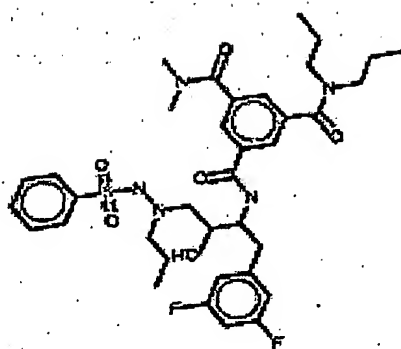
1B4 1,1,4,4



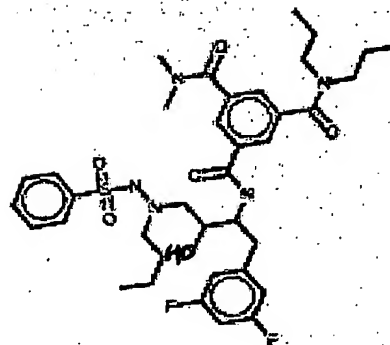
1B5 1,2,1,1



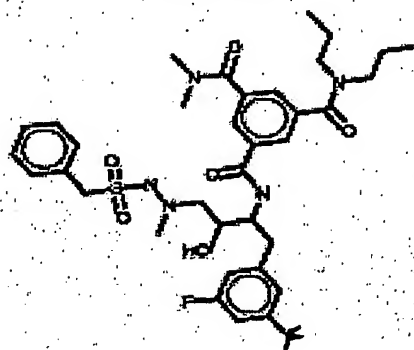
1B6 1,2,1,2



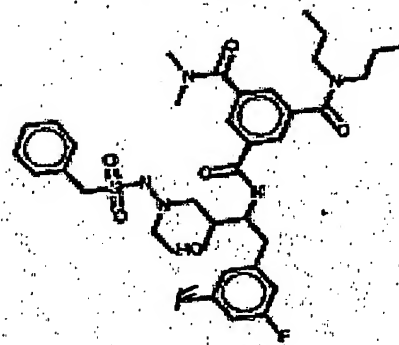
1:87 1.2.1.3



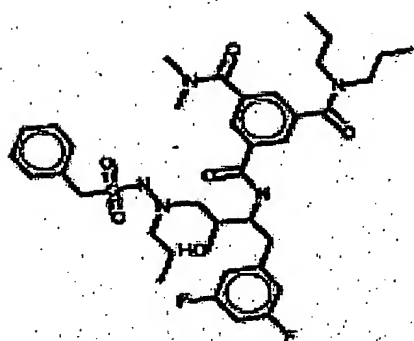
1:88 1.2.1.4



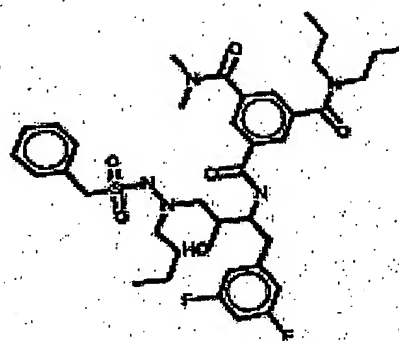
1:89 1.2.2.1



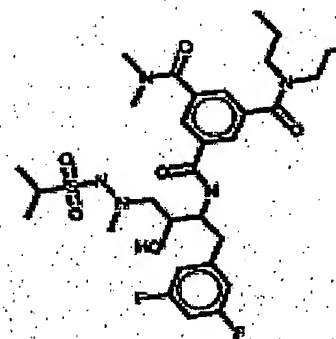
1:90 1.2.2.2



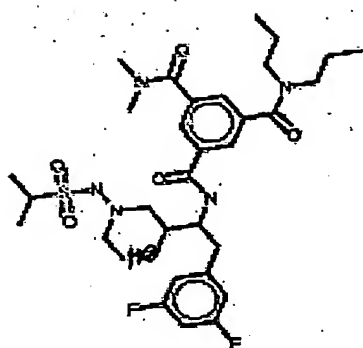
1:91 1.2.2.3



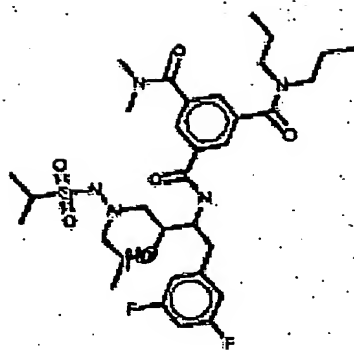
1:912 1.2.2.4



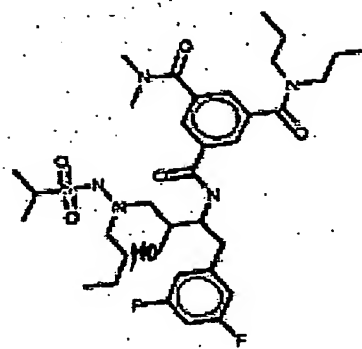
1:91 1.2.3.1



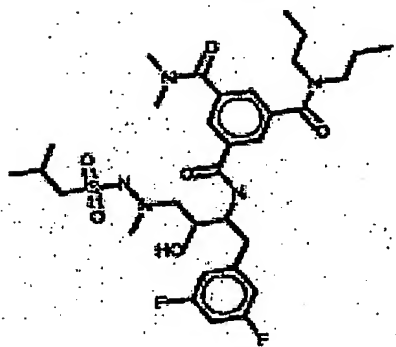
1C2 1,2,3,2



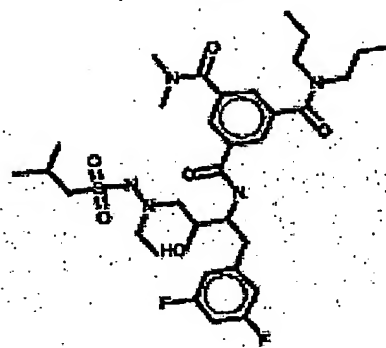
1C3 1,2,3,3



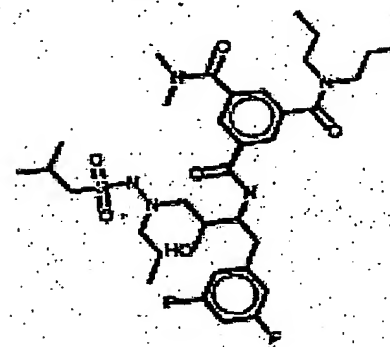
1C4 1,2,3,4



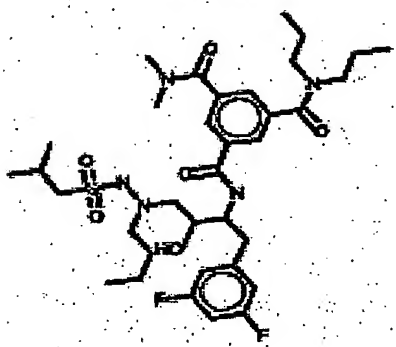
1C5 1,2,4,1



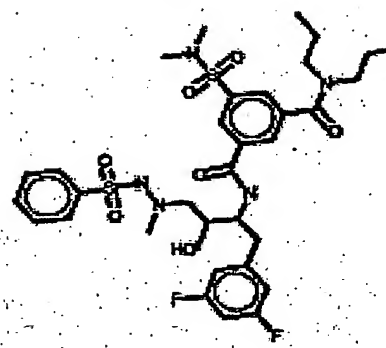
1C6 1,2,4,2



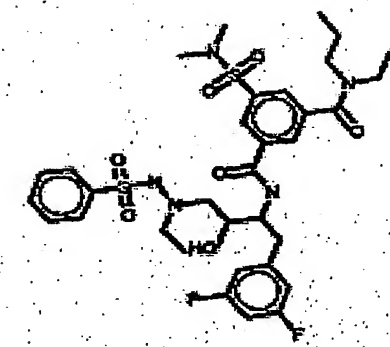
1C7 1,2,4,3



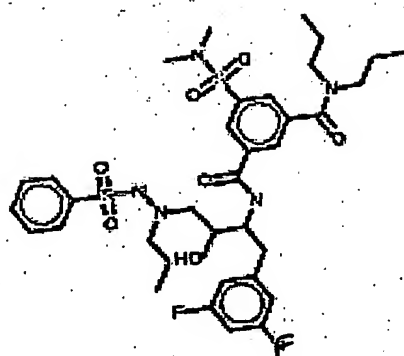
1C8 1,2,4,4



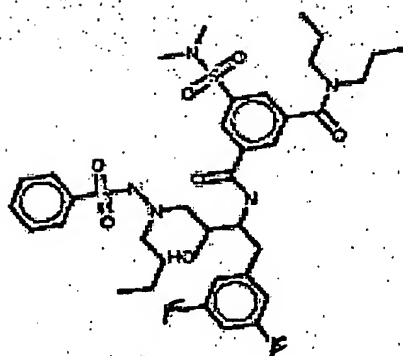
1C9 1,3,1,1



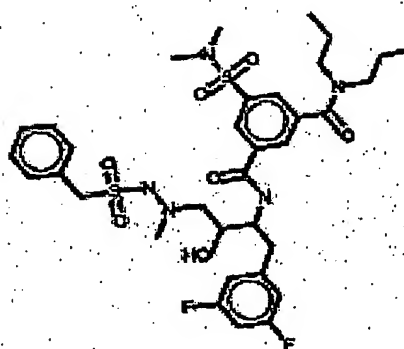
1C10 1,3,1,2



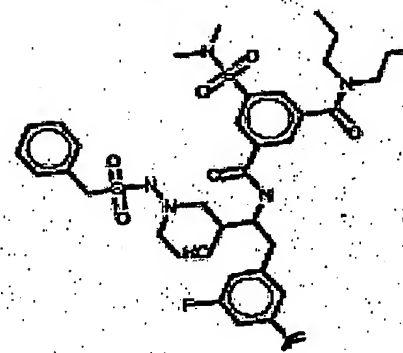
1:G11 1,3,1,3



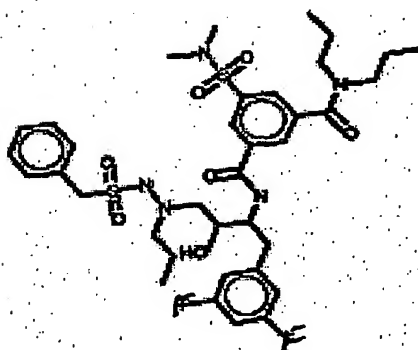
1:G12 1,3,1,4



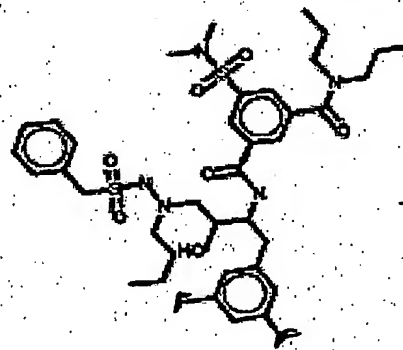
1:D1 1,3,2,1



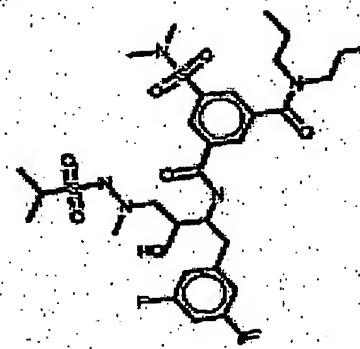
1:D2 1,3,2,2



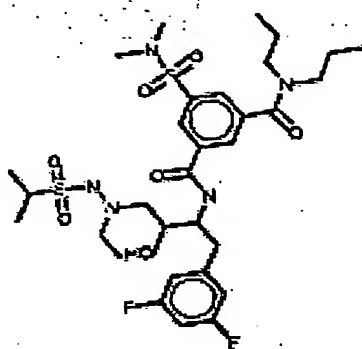
1:G9 1,3,2,3



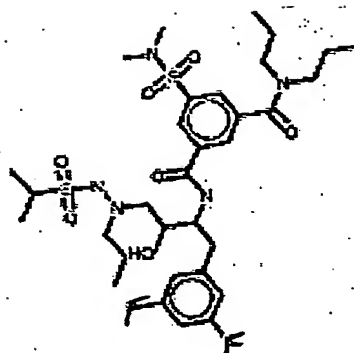
1:D4 1,3,2,4



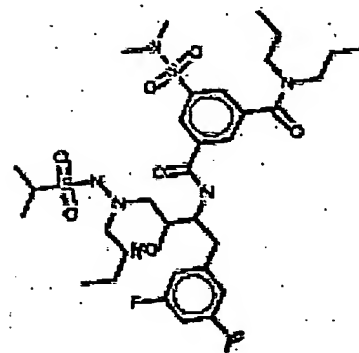
1:G6 1,3,3,1



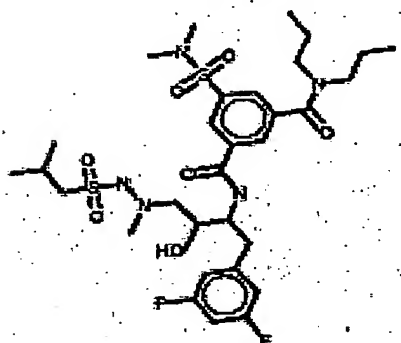
1:06 1,3,3,2



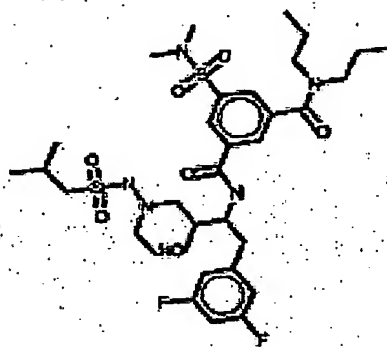
1:07 1,3,3,3



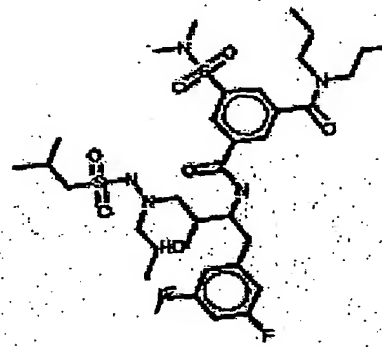
1:08 1,3,3,4



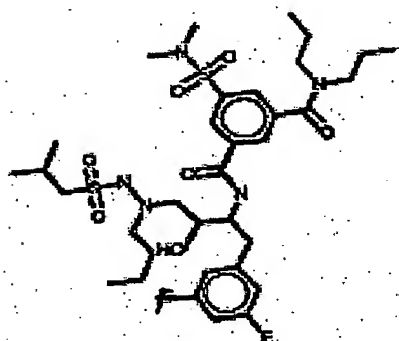
1:09 1,3,4,1



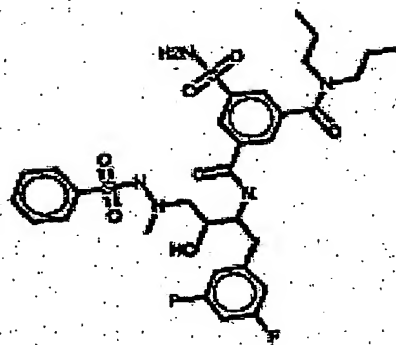
1:10 1,3,4,2



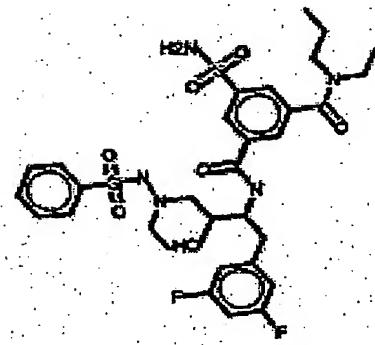
1:11 1,3,4,3



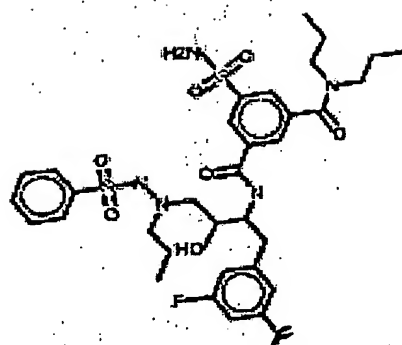
1:12 1,3,4,4



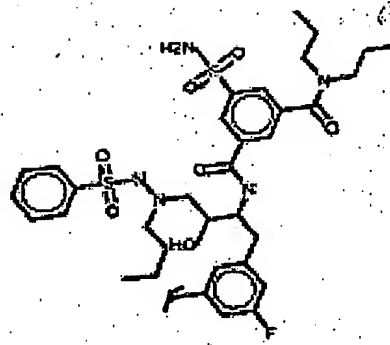
1:11 1,4,1,1



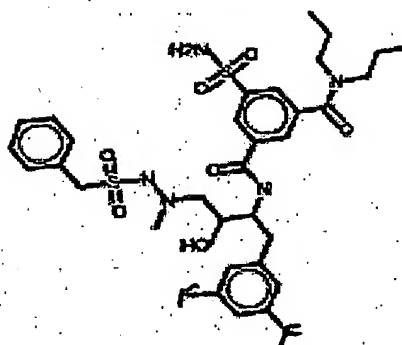
1:12 1,4,1,2



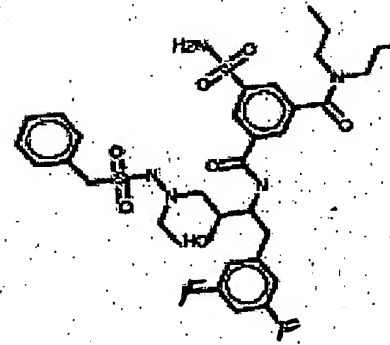
1:ES 1,4,1,3



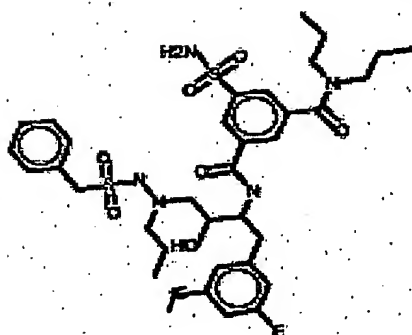
1:ES 1,4,1,4



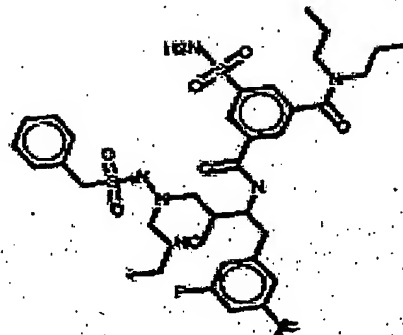
1:ES 1,4,2,1



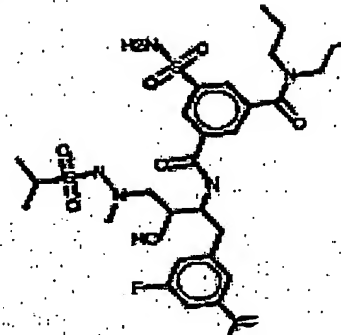
1:ES 1,4,2,2



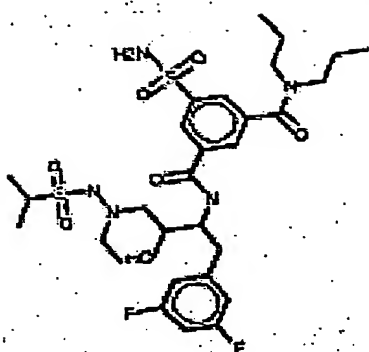
1:ES 1,4,2,3



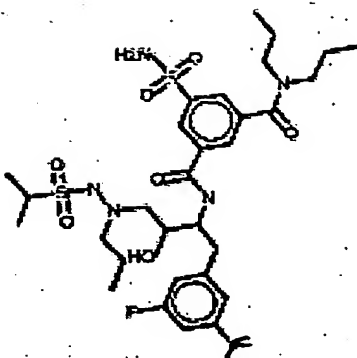
1:ES 1,4,2,4



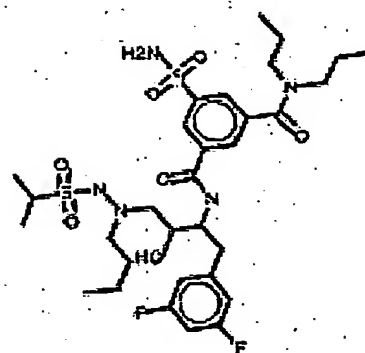
1:ES 1,4,2,1



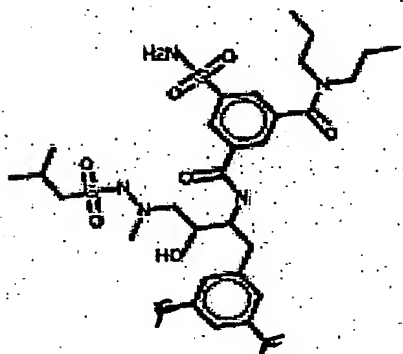
1-E10 1,4,3,2



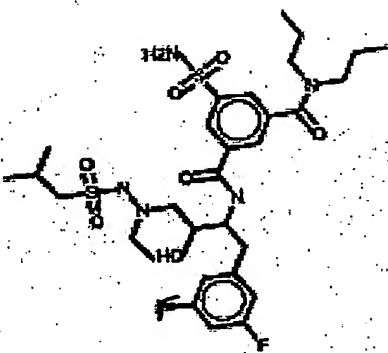
1-E11 1,4,3,3



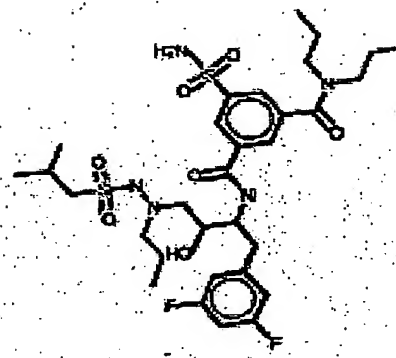
1-E12 1,4,3,4



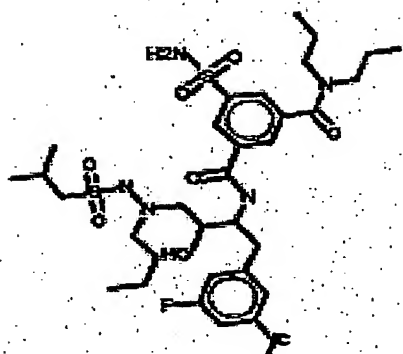
1-F1 1,4,4,1



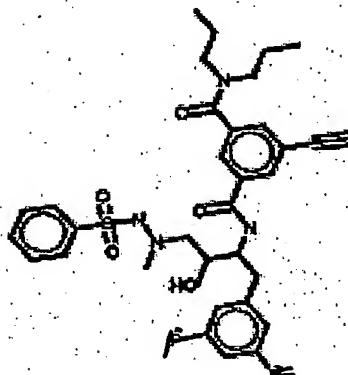
1-F2 1,4,4,2



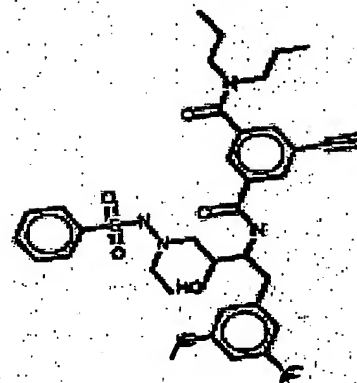
1-F3 1,4,4,3



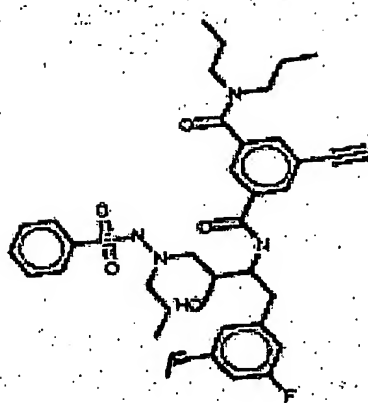
1-F4 1,4,4,4



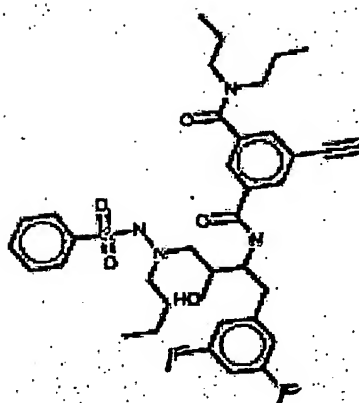
1-F5 1,5,1,1



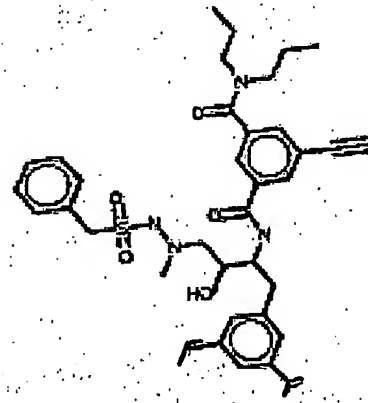
1-F6 1,5,1,2



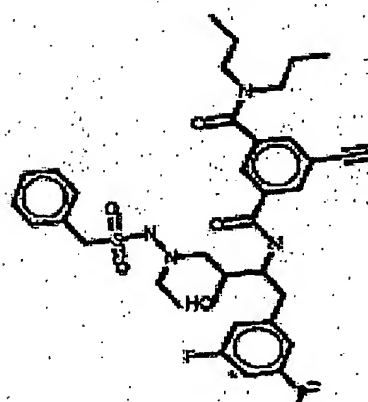
1F7 1,5,1,3



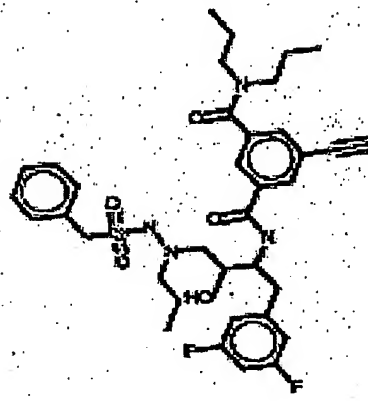
1F8 1,5,1,4



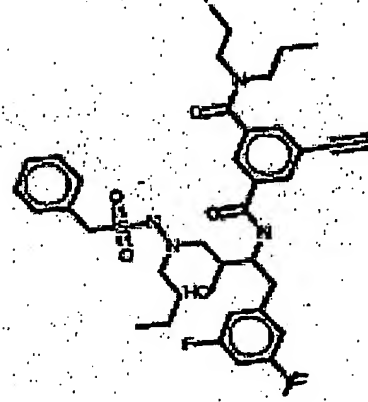
1F9 1,5,2,1



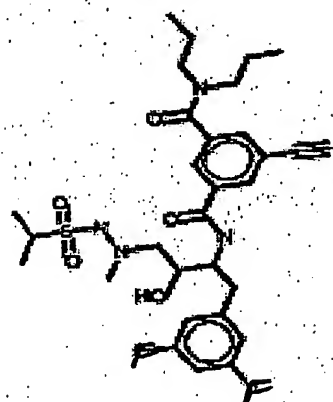
1F10 1,5,2,2



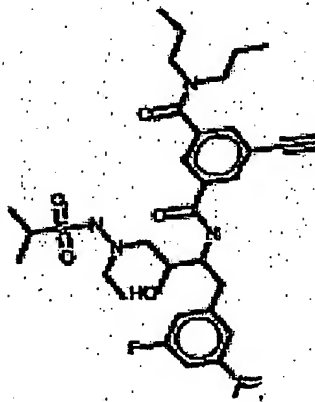
1F11 1,5,2,3



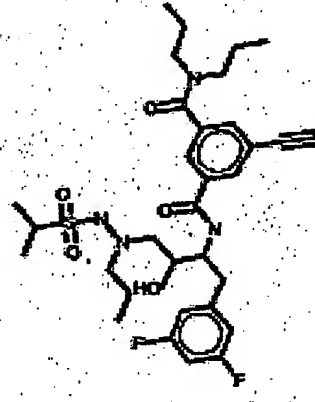
1F12 1,5,2,4



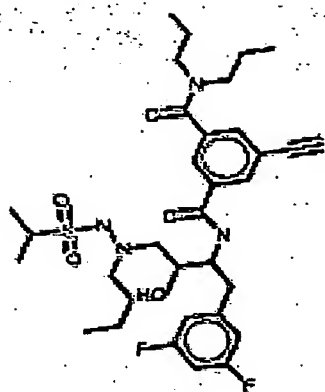
1G1 1,5,3,1



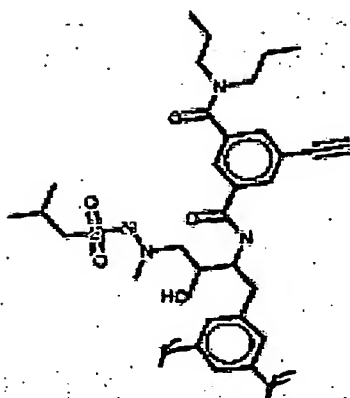
1G2 1,5,3,2



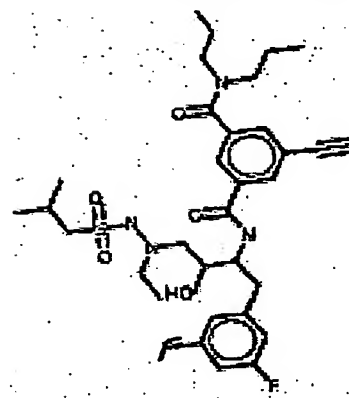
1G3 1,5,3,3



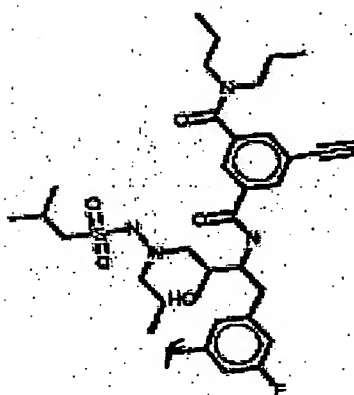
1:G4 1,5,3,4



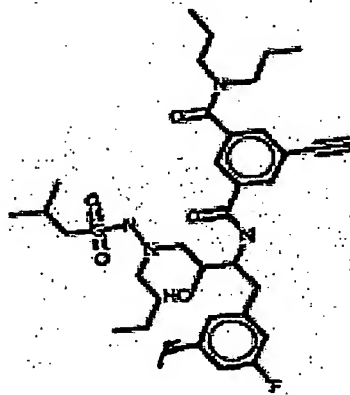
1:G5 1,5,4,1



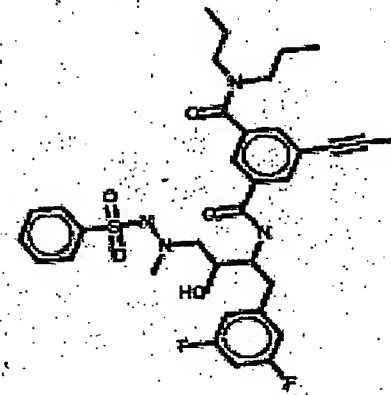
1:G6 1,5,4,2



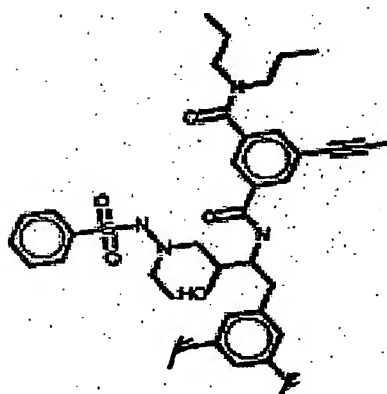
1:G7 1,5,4,3



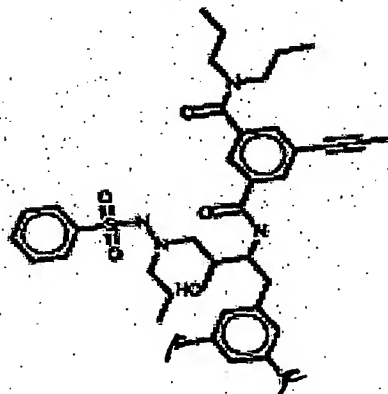
1:G8 1,5,4,4



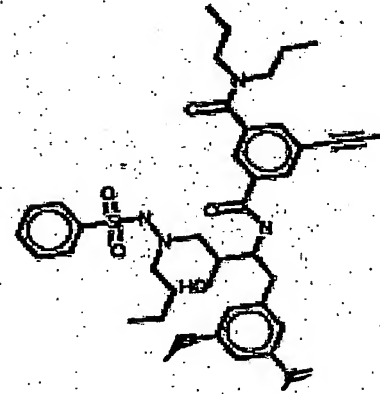
1:G9 1,6,1,1



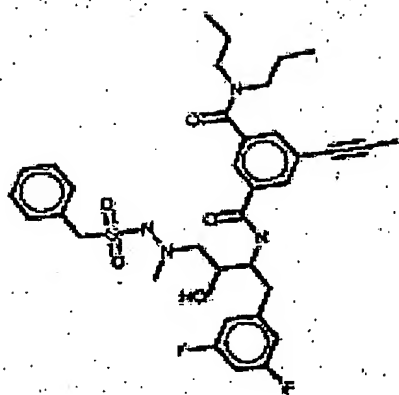
1:G10 1,6,1,2



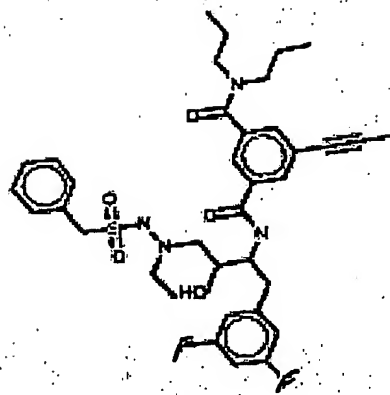
1:G11 1,6,1,3



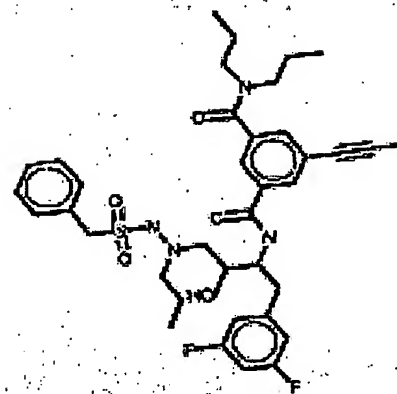
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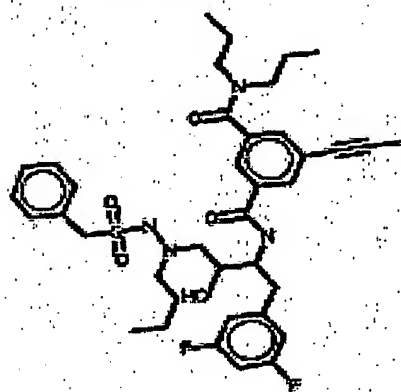
1111 1.6.2.1



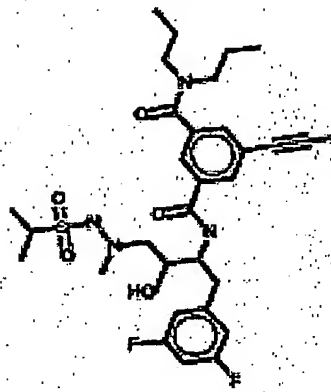
1112 1.6.2.2



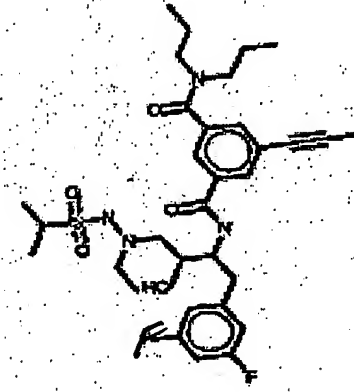
1113 1.6.2.3



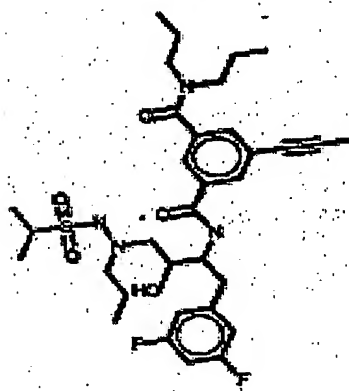
1114 1.6.2.4



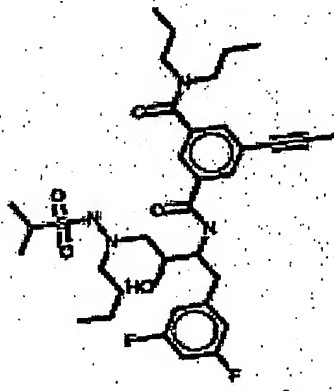
1115 1.6.2.1



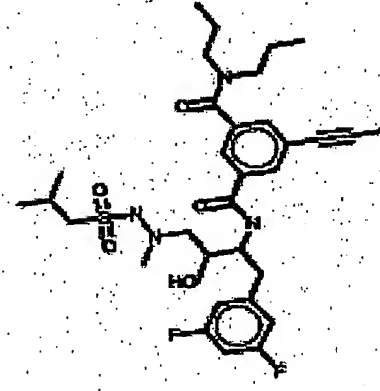
1116 1.6.2.2



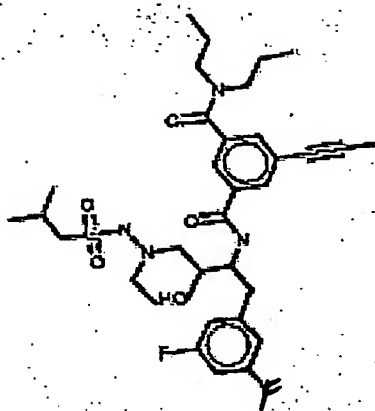
1117 1.6.2.3



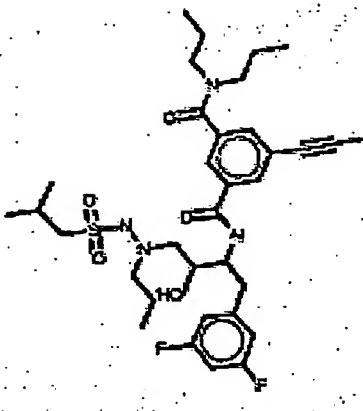
1118 1.6.2.4



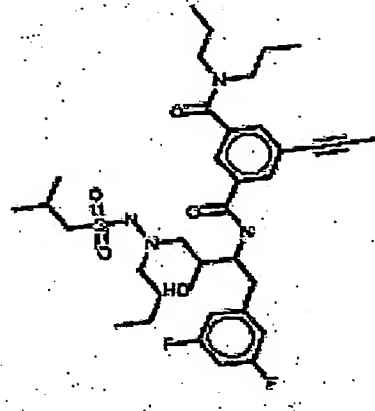
1119 1.6.4.1



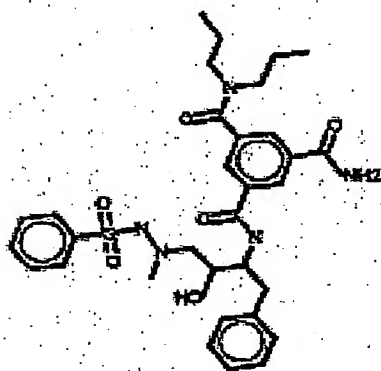
1A42



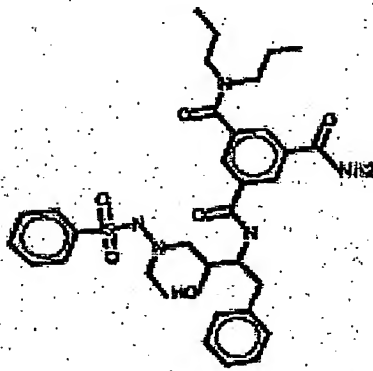
1A43



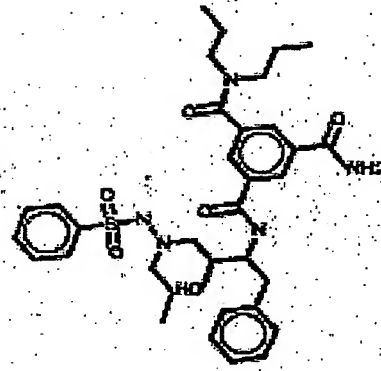
1A44



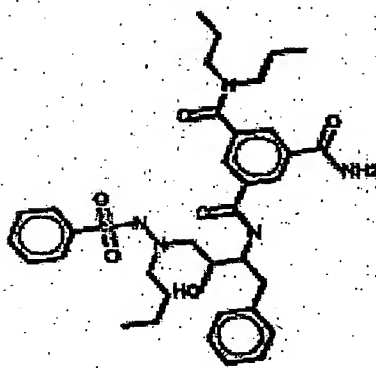
2A1



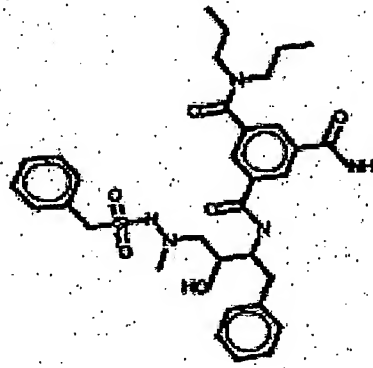
2A2



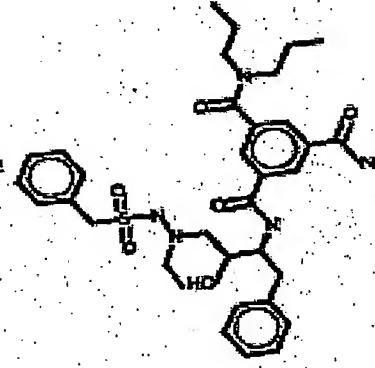
2A3



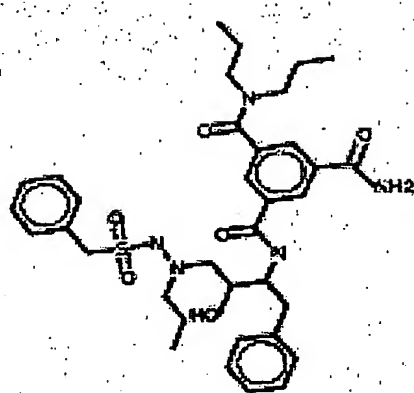
2A4



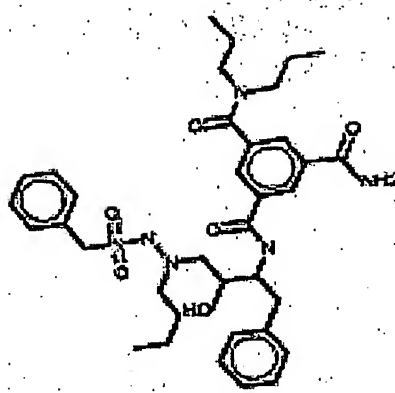
2A5



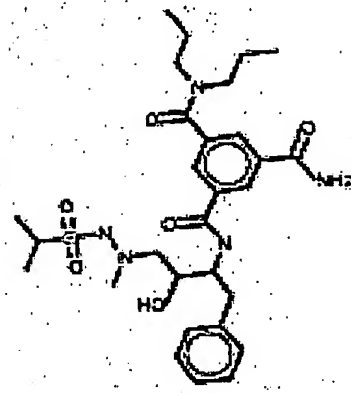
2A6



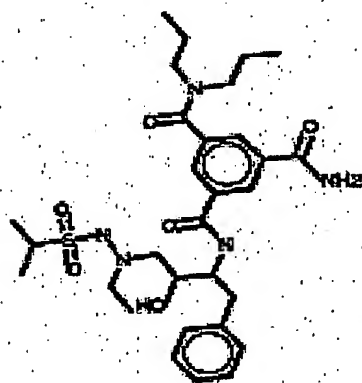
2A7 2,1,2,3



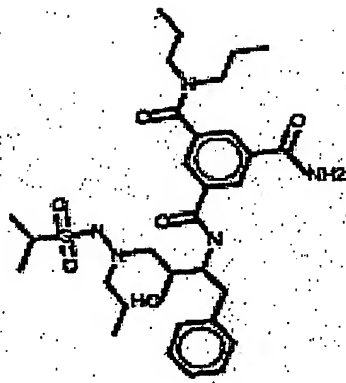
2A8 2,1,2,4



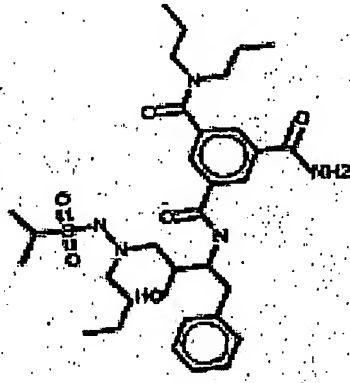
2A9 2,1,3,1



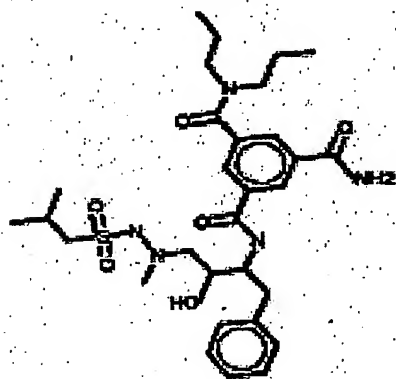
2A10 2,1,3,2



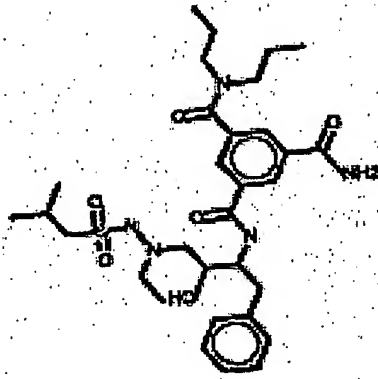
2A11 2,1,3,3



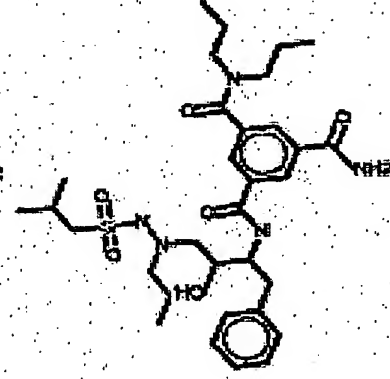
2A12 2,1,3,4



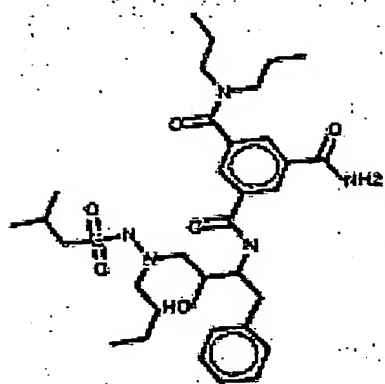
2B1 2,1,4,1



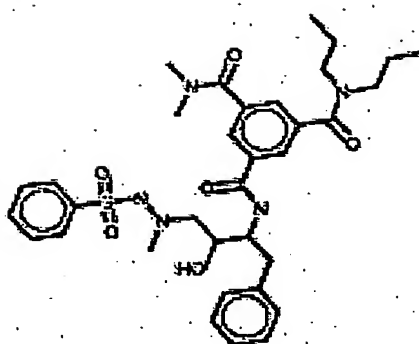
2B2 2,1,4,2



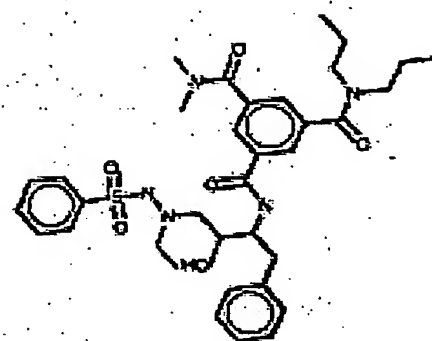
2B3 2,1,4,3



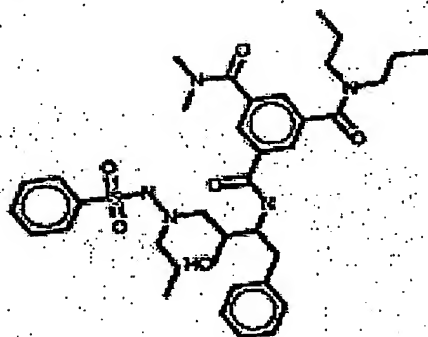
284 2.1.4.4



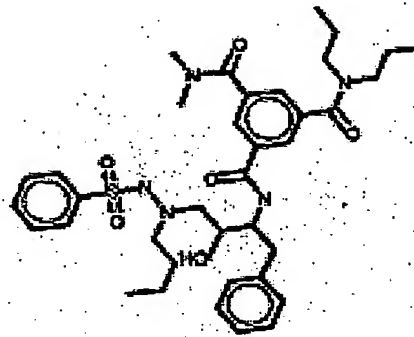
285 2.2.1.1



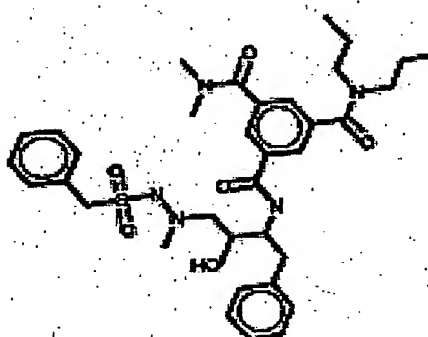
286 2.2.1.2



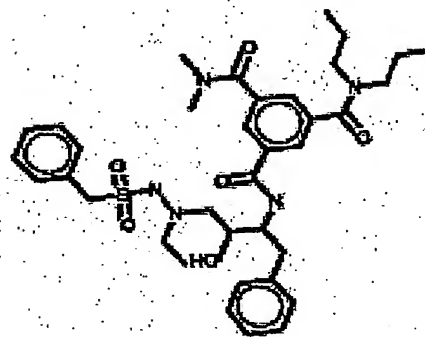
287 2.2.1.3



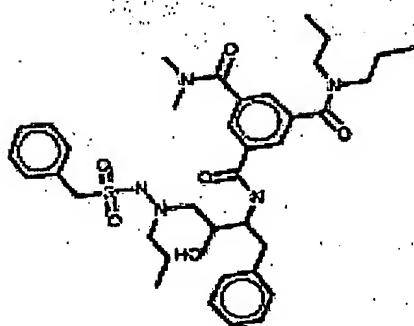
288 2.2.1.4



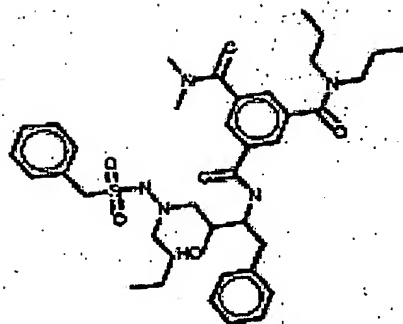
289 2.2.2.1



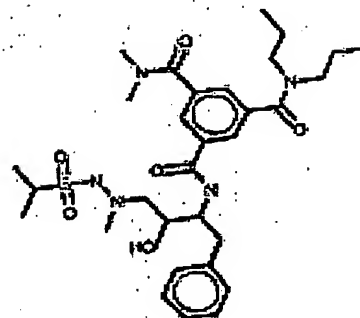
290 2.2.2.2



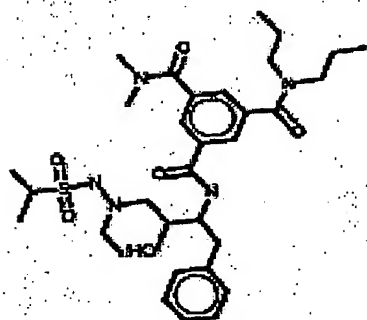
2C11 2.2.2.3



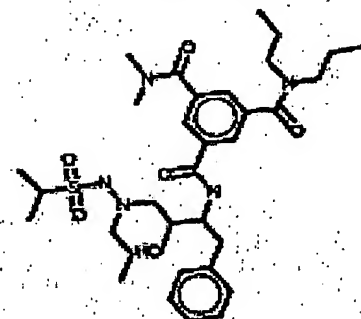
2C12 2.2.2.4



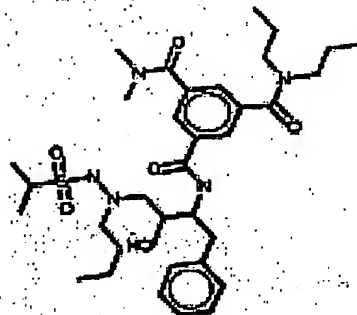
2C1 2.2.3.1



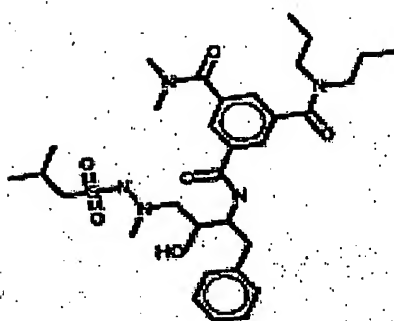
2C2 2.2.3.2



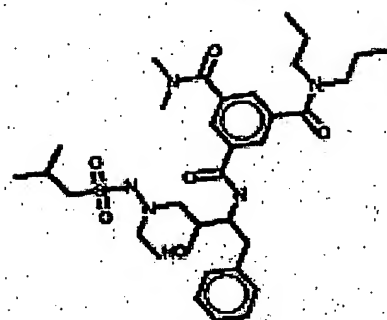
2C3 2.2.3.3



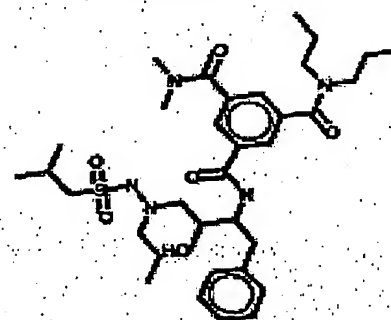
2C4 2.2.3.4



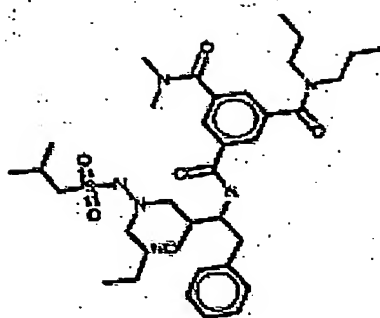
2C5 2.2.4.1



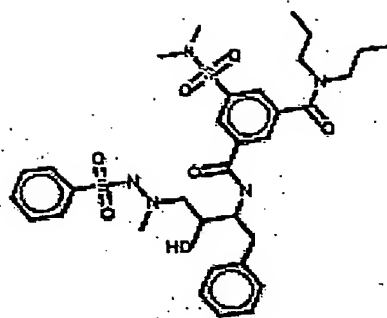
2C6 2.2.4.2



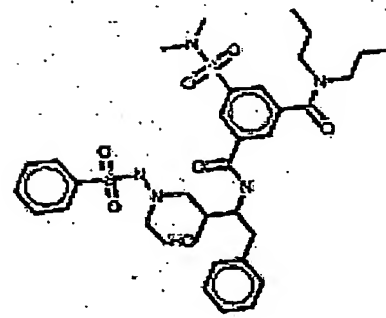
2C7 2.2.4.3



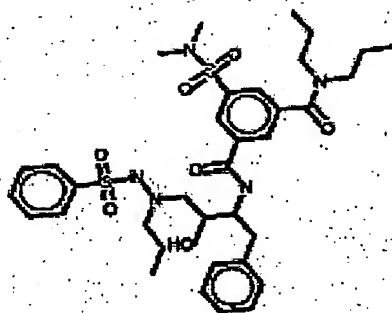
2C8 2.2.4.4



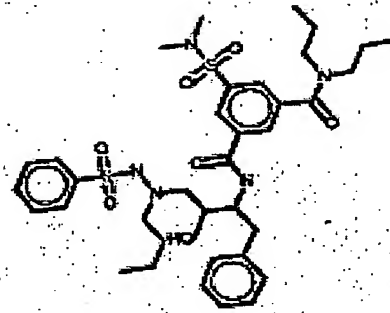
2C9 2.3.1.1



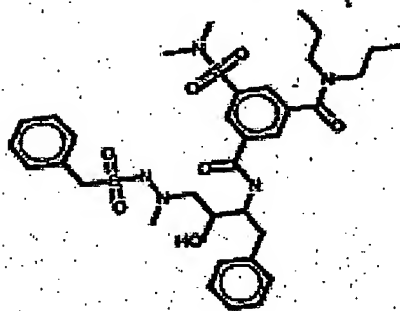
2C10 2.3.1.2



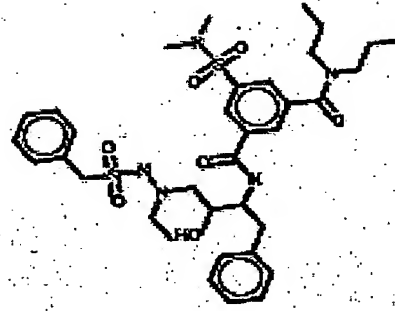
2C11 2.3.1.3



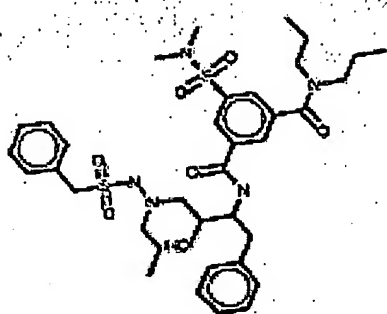
2C12 2.3.1.4



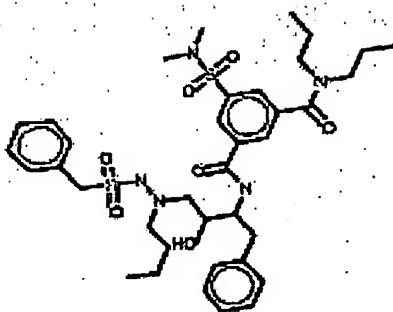
2C1 2.3.2.1



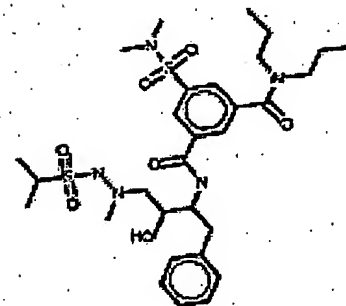
2C2 2.3.2.2



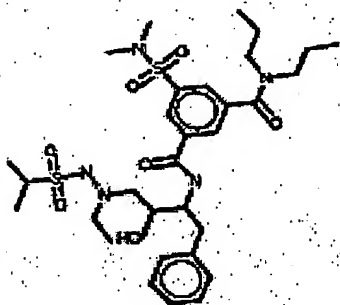
2D3 2.3.2.3



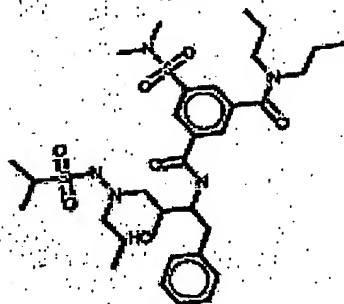
2D4 2.3.2.4



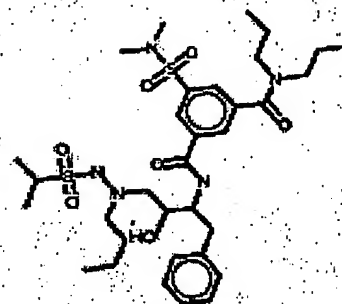
2D5 2.3.3.1



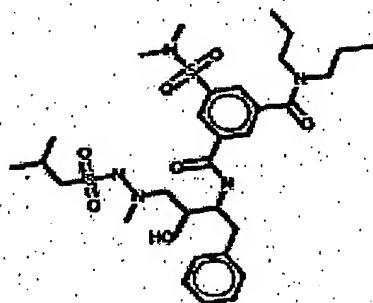
2D6 2.3.3.2



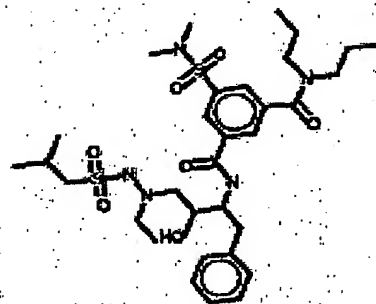
2D7 2.3.3.3



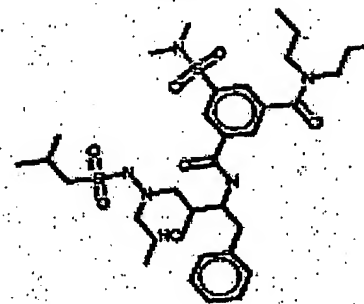
2D8 2.3.3.4



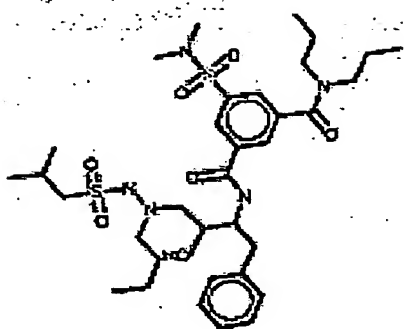
2D9 2.3.4.1



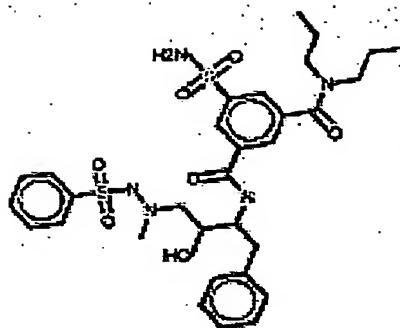
2D10 2.3.4.2



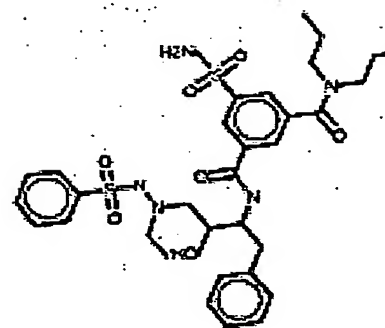
2D11 2.3.4.3



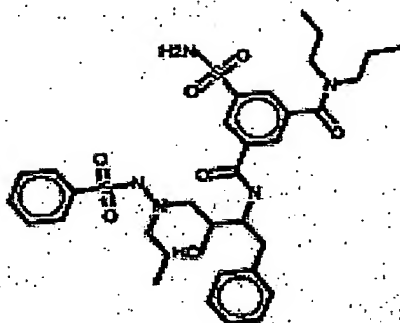
2D12 2,3,4,4



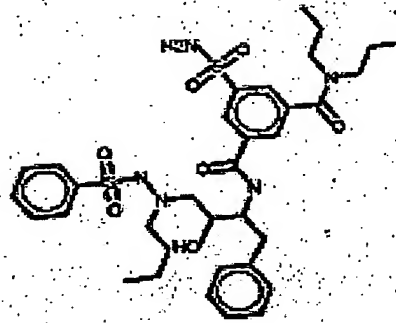
2E1 2,4,3,1



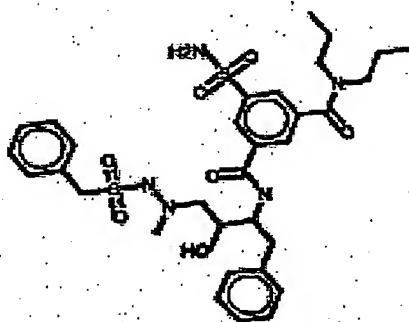
2E2 2,4,1,3



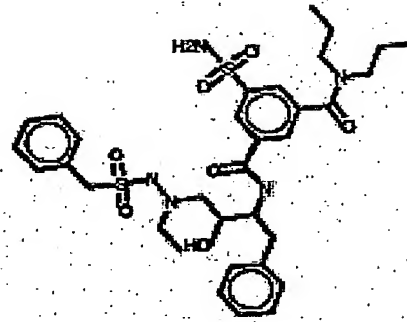
2E3 2,4,1,3



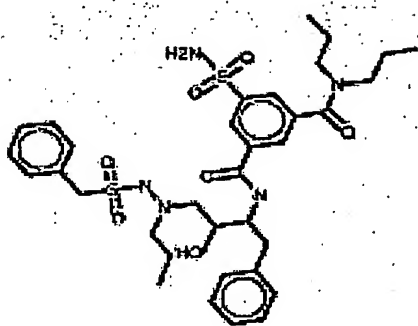
2E4 2,4,1,4



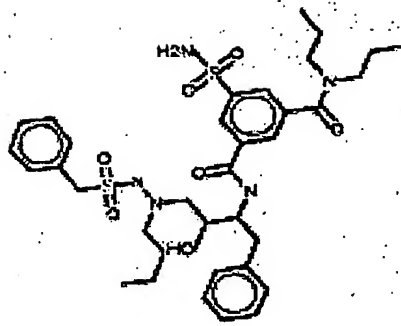
2E5 2,4,2,1



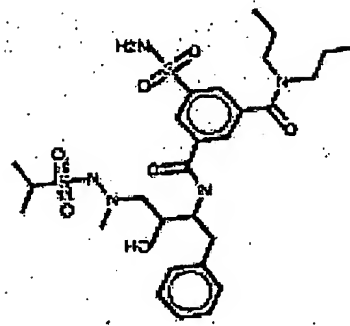
2E6 2,4,2,2



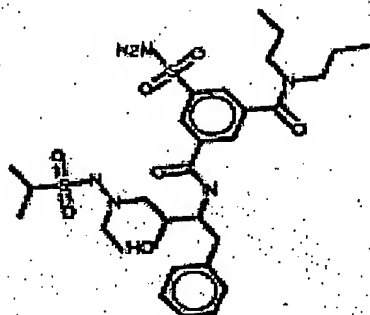
2E7 2,4,2,3



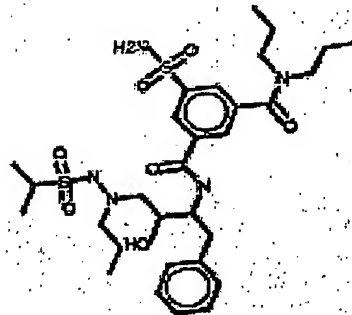
2E8 2,4,2,4



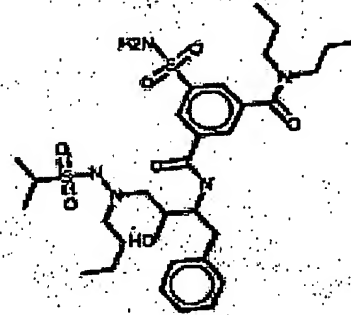
2E9 2,4,3,1



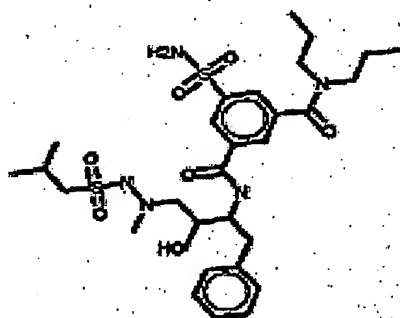
2E10 2,4,3,2



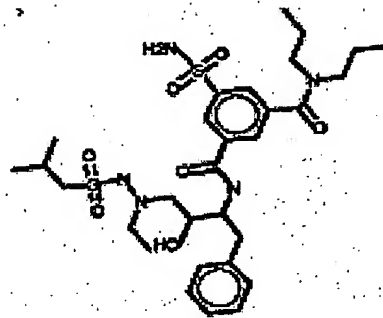
2E11 2,4,3,3



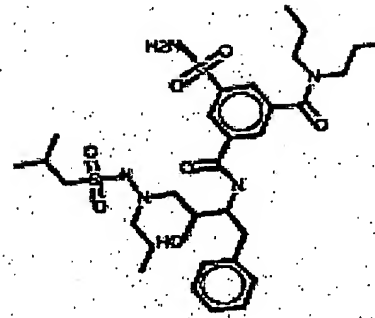
2E12 2,4,3,4



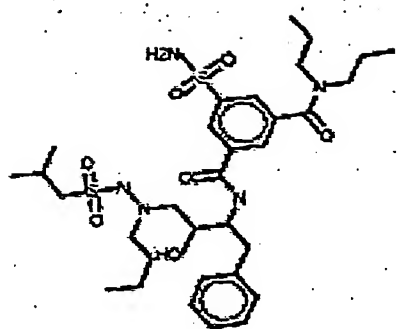
2F1 2,4,4,1



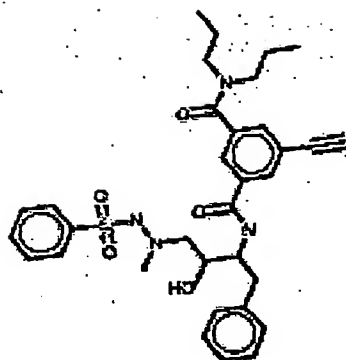
2F2 2,4,4,2



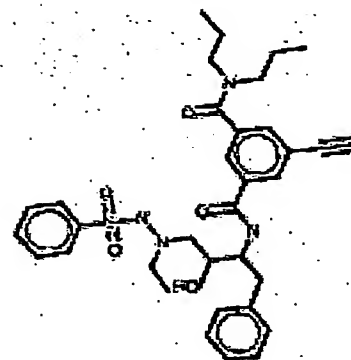
2F3 2,4,4,3



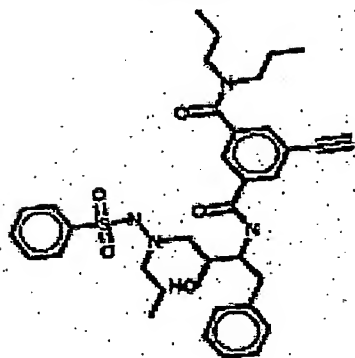
2F4 2,4,4,4



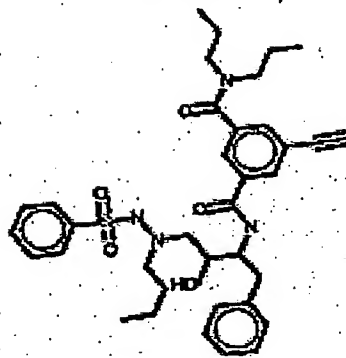
2F5 2,5,1,1



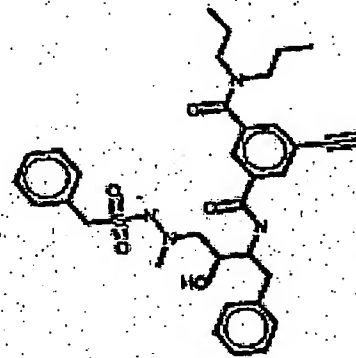
2F6 2,5,1,2



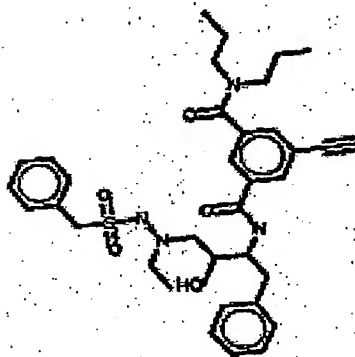
2F7 2,5,1,3



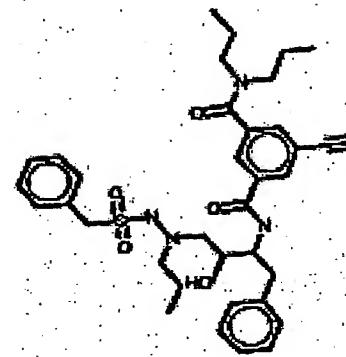
2F8 2,5,1,4



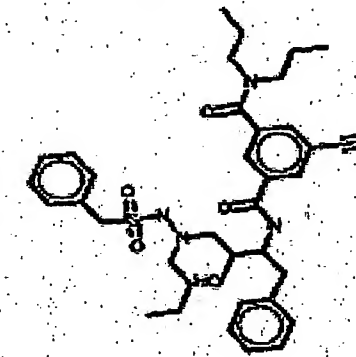
2F9 2,5,2,1



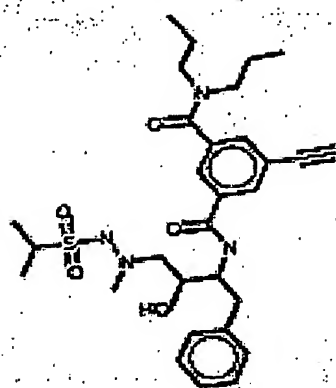
2F10 2,5,2,2



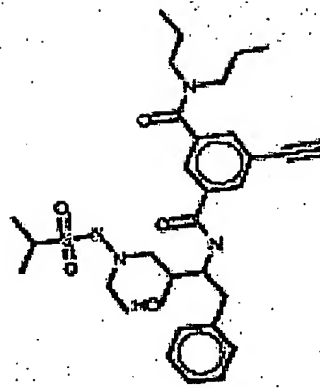
2F11 2,5,2,3



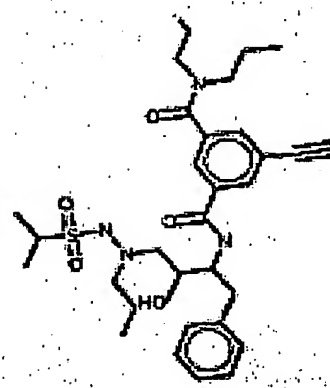
2F12 2,5,2,4



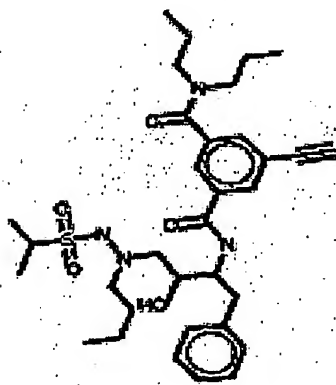
2G1 2.5.3.1



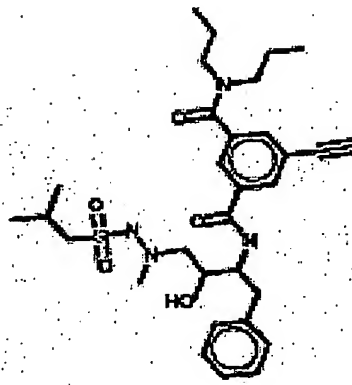
2G2 2.5.3.2



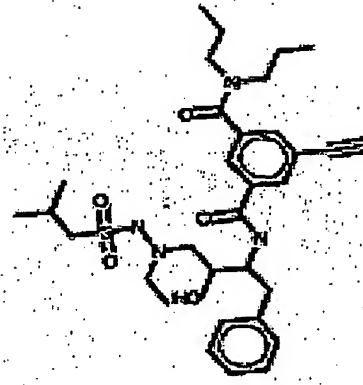
2G3 2.5.3.3



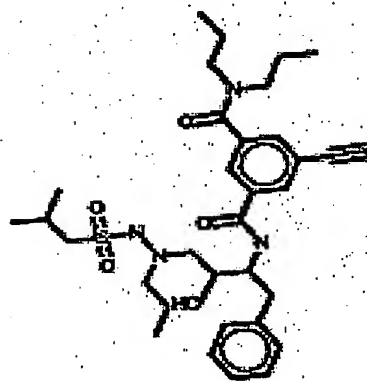
2G4 2.5.3.4



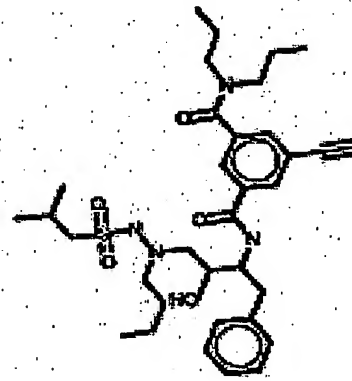
2G5 2.5.4.1



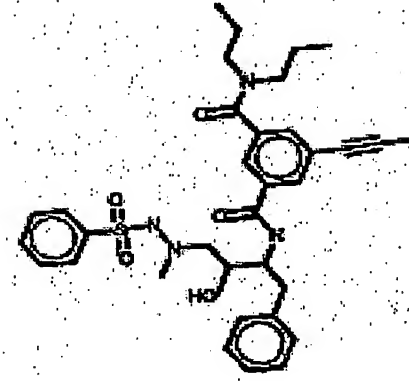
2G6 2.5.4.2



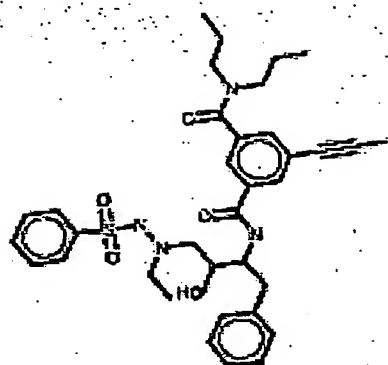
2G7 2.5.4.3



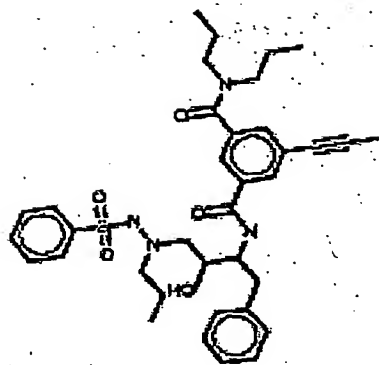
2G8 2.5.4.4



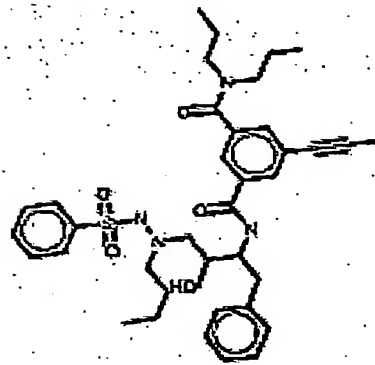
2G9 2.5.4.5



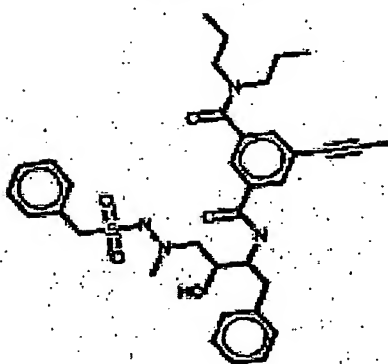
2G10 2.6.1.2



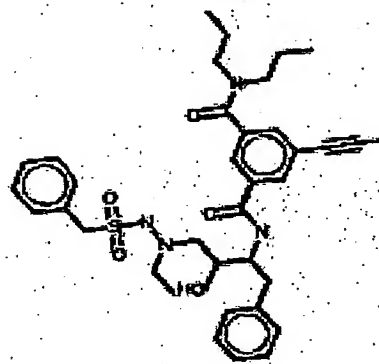
2G11 2.6.1.3



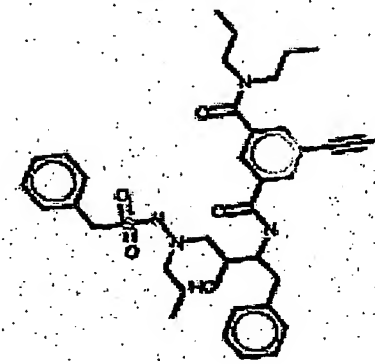
2G12 2.6.1.4



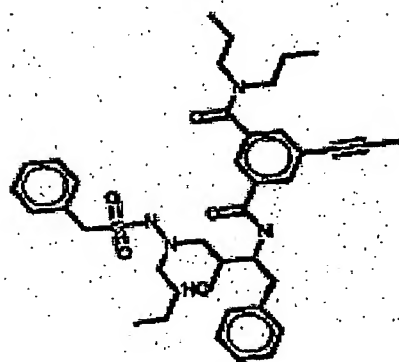
2H1 2.6.2.1



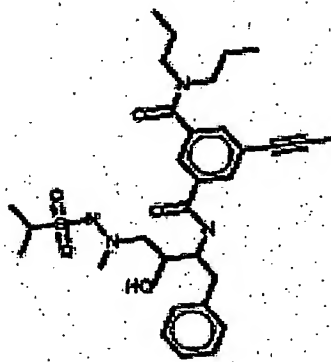
2H2 2.6.2.2



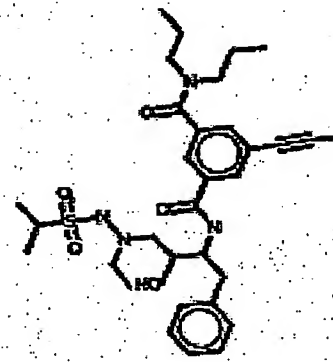
2H3 2.6.2.3



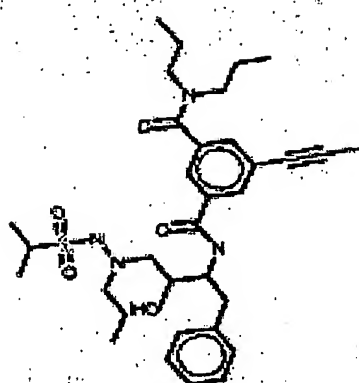
2H4 2.6.2.4



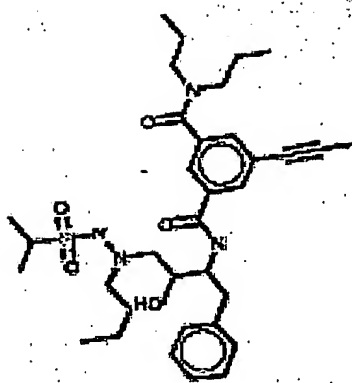
2H5 2.6.3.1



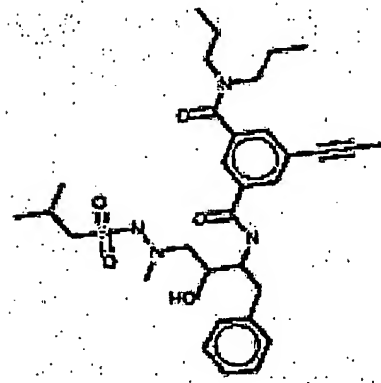
2H6 2.6.3.2



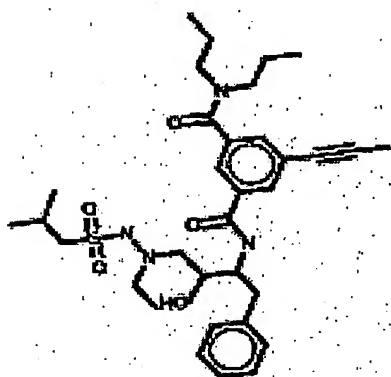
2H7 2.6.3.3



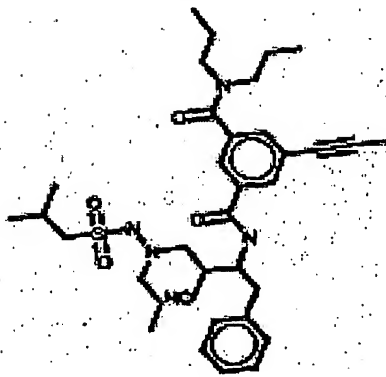
2H8 2.6.3.4



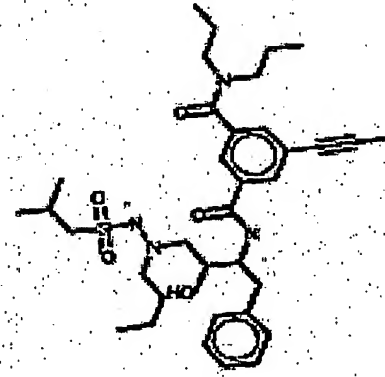
2H9 2.6.4.1



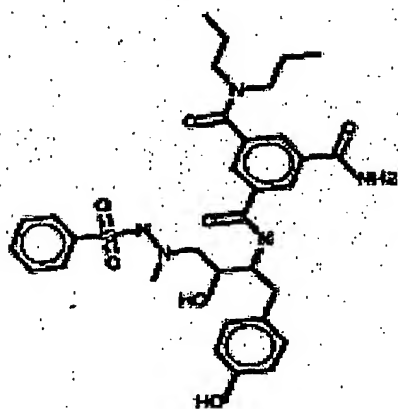
2H10 2.6.4.2



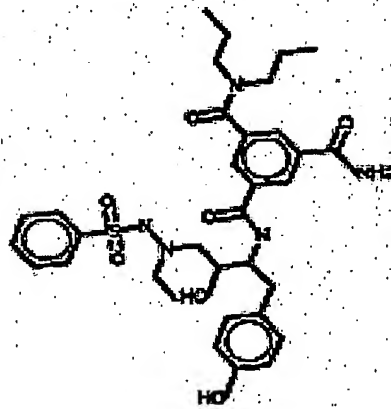
2H11 2.6.4.3



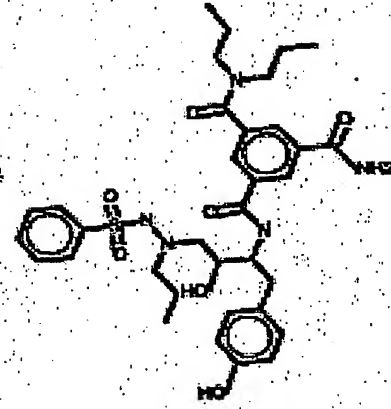
2H12 2.6.4.4



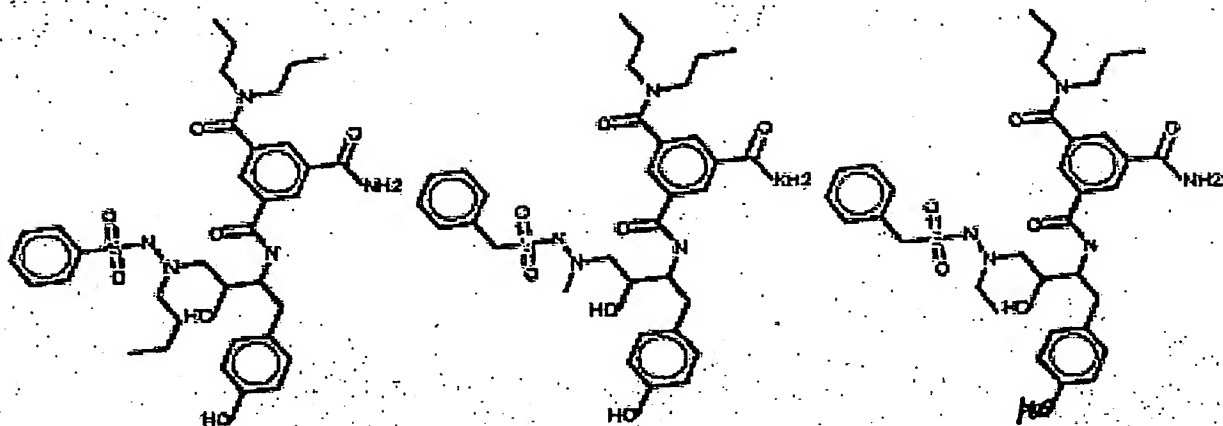
3A1 3.1.1.1



3A2 3.1.1.2



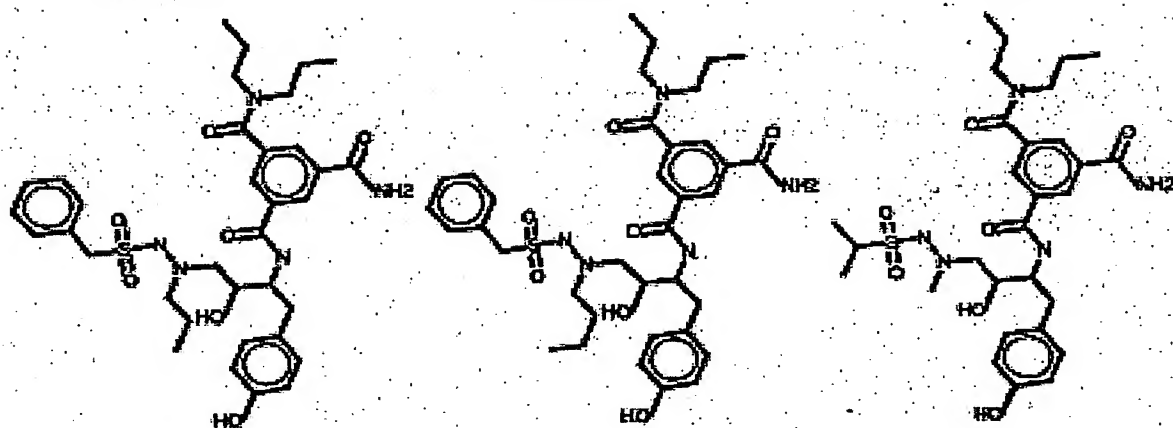
3A3 3.1.1.3



3A4 3,1,1,4

3A5 3,1,2,1

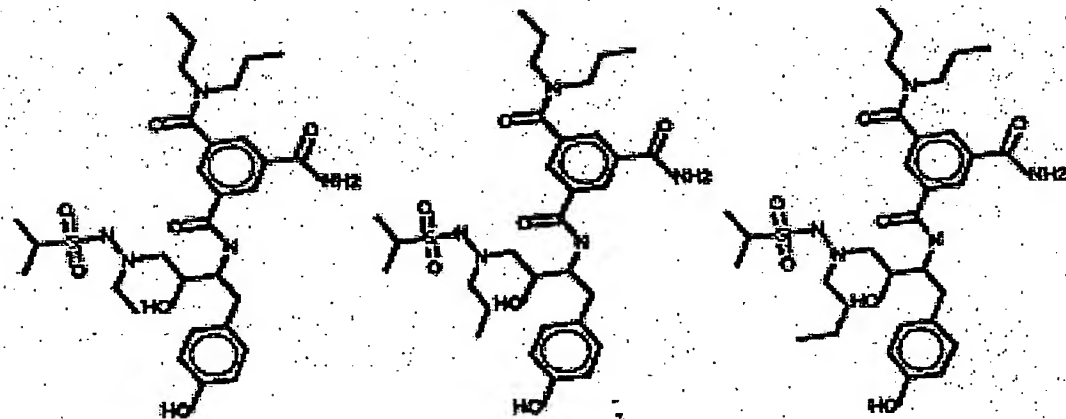
3A6 1,1,2,2



3A7 3,1,2,3

3A8 3,1,2,4

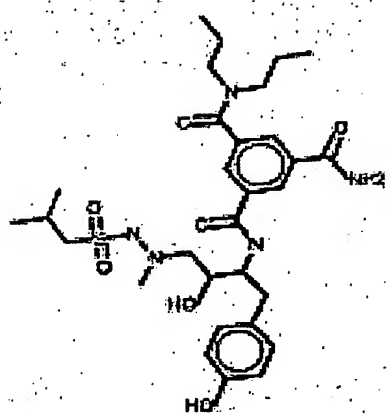
3A9 3,1,3,1



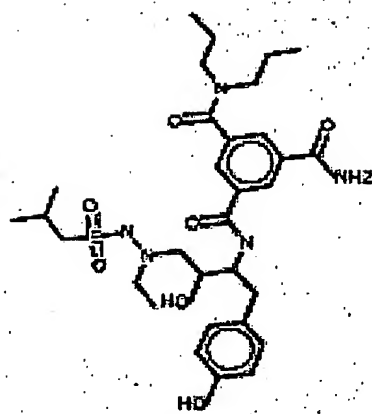
3A10 3,1,3,2

3A11 3,1,3,3

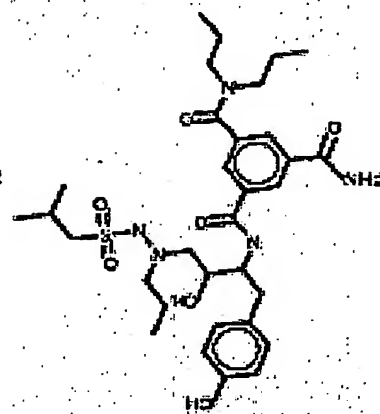
3A12 3,1,3,4



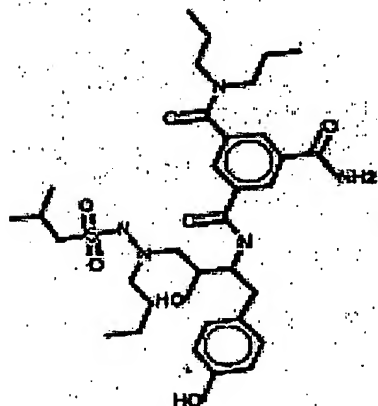
3B1 3.1.4.1



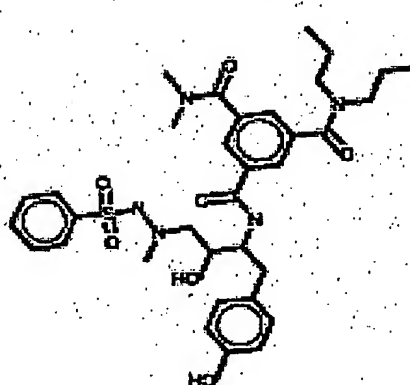
3B2 3.1.4.2



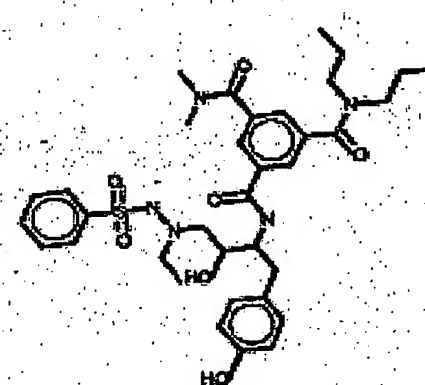
3B3 3.1.4.3



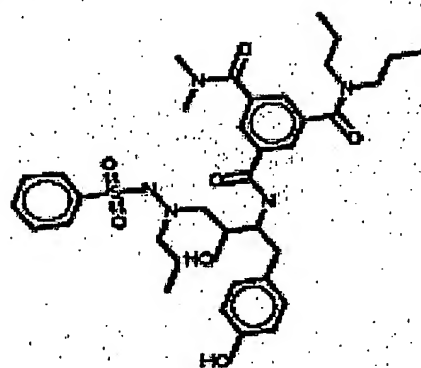
3B4 3.1.4.4



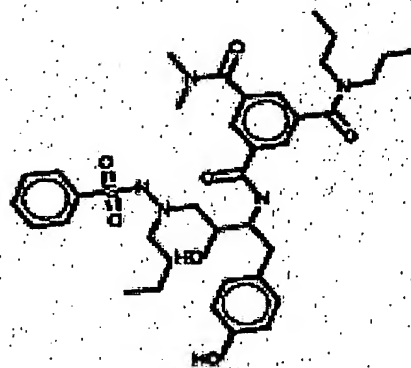
3B5 3.2.1.1



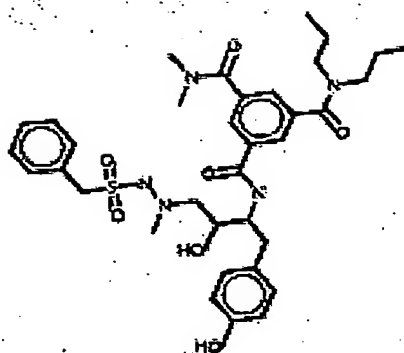
3B6 3.2.1.2



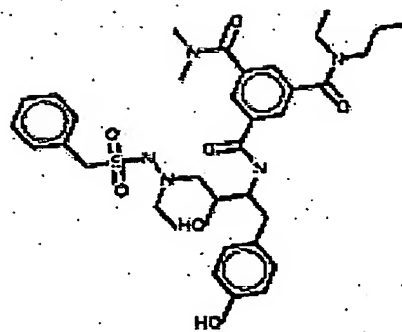
3B7 3.2.1.3



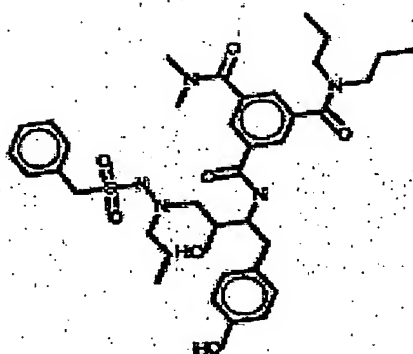
3B8 3.2.1.4



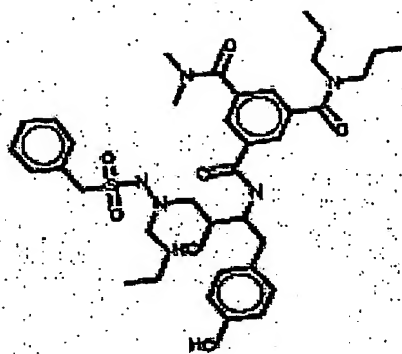
3B9 3.2.2.1



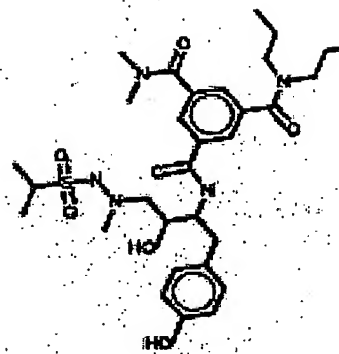
3B10 3.2.2.2



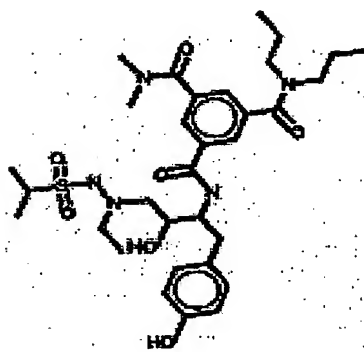
3B11 3.2.2.3



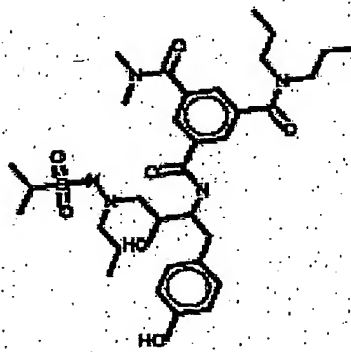
3B12 3.2.2.4



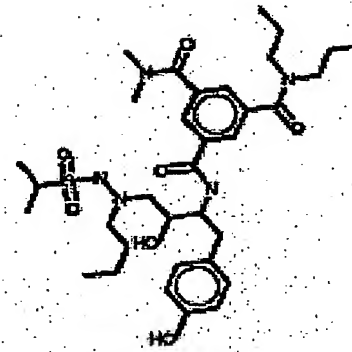
3C1 3.2.3.1



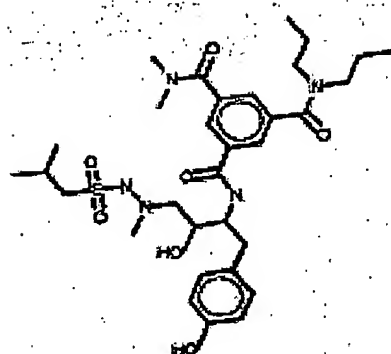
3C2 3.2.3.2



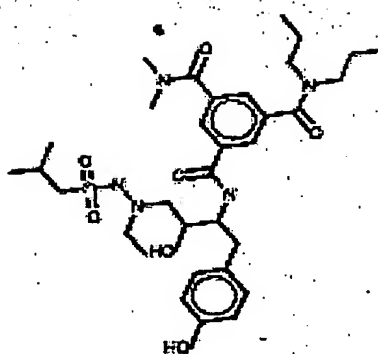
3C3 3.2.3.3



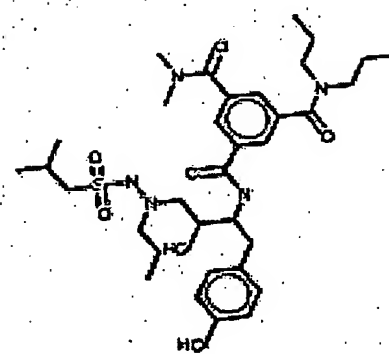
3C4 3.2.3.4



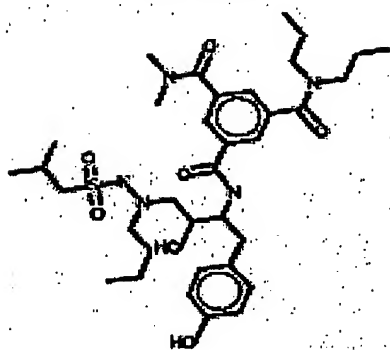
3C8 3.2.4.1



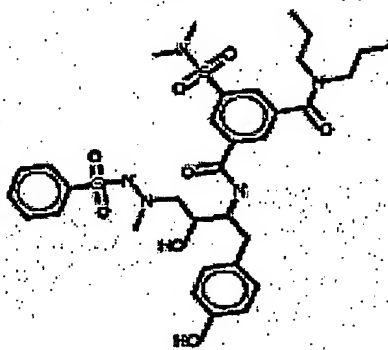
3C8 3.2.4.2



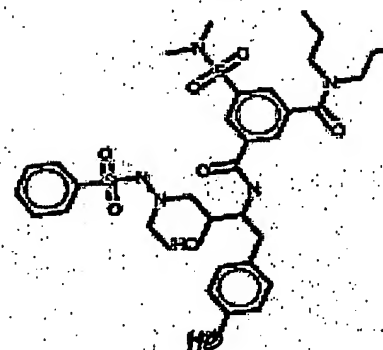
3C7 3.2.4.3



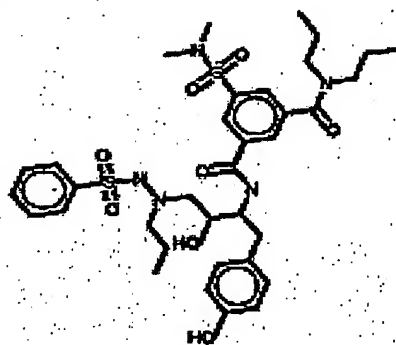
3C8 3.2.4.4



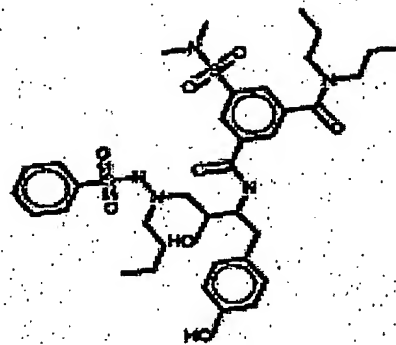
3C9 3.3.1.1



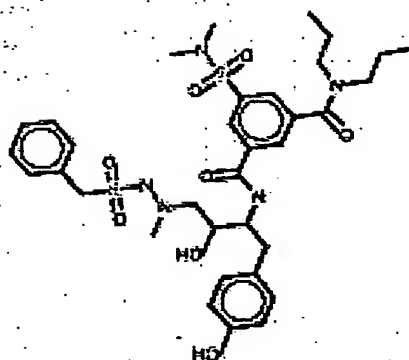
3C10 3.3.1.2



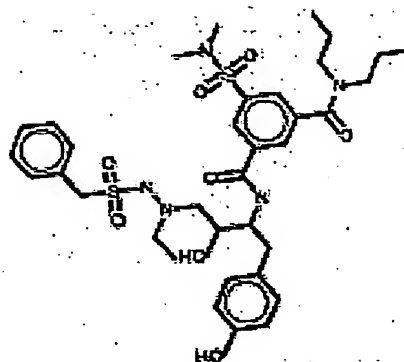
3C11 3.3.1.3



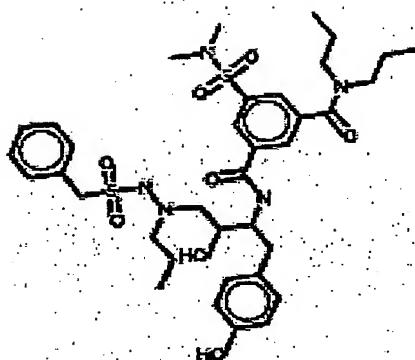
3C12 3.3.1.4



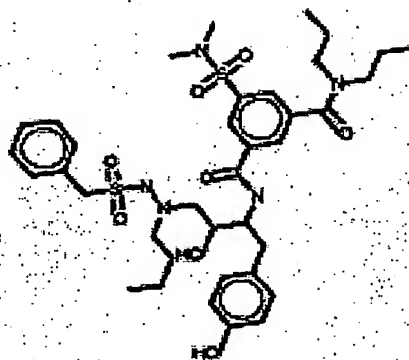
301 3.3.2.1



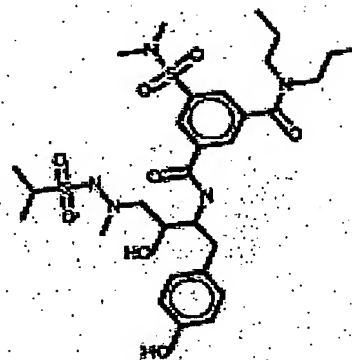
302 3.3.2.2



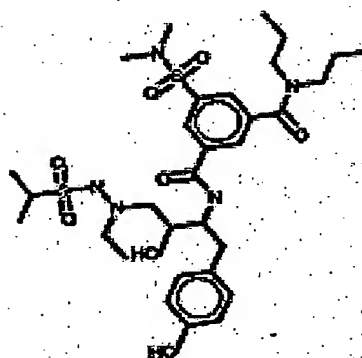
303 3.3.2.3



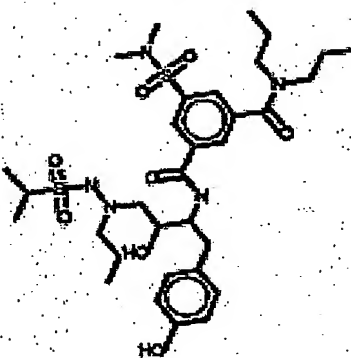
304 3.3.2.4



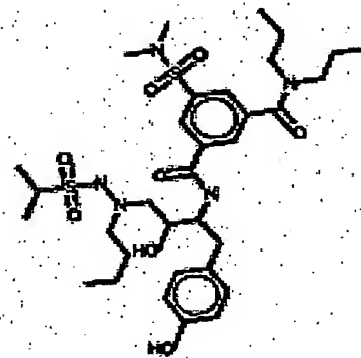
305 3.3.3.1



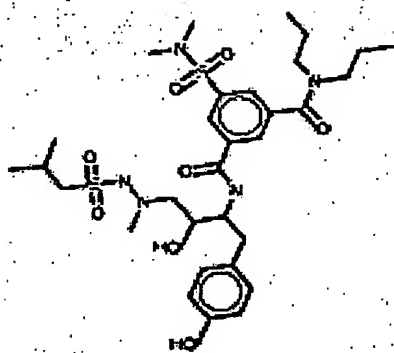
306 3.3.3.2



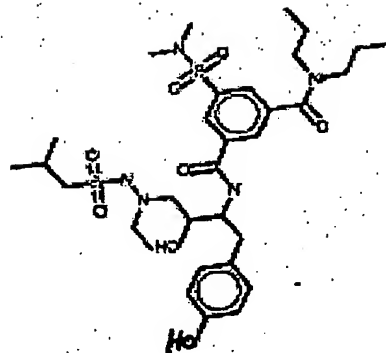
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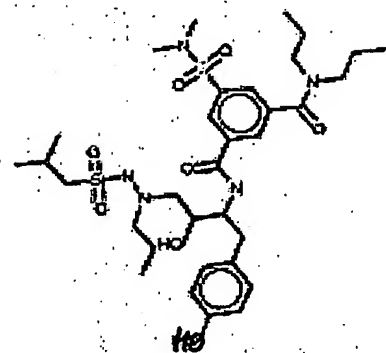
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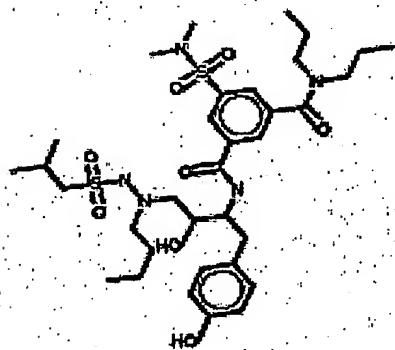
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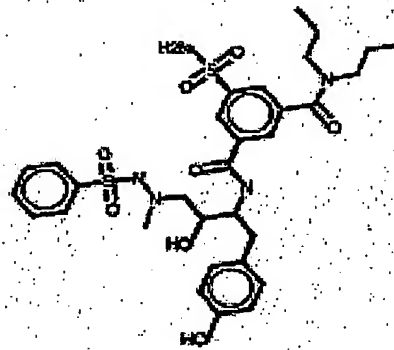
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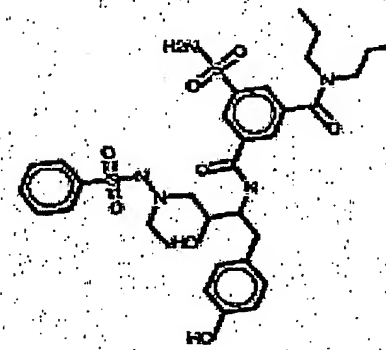
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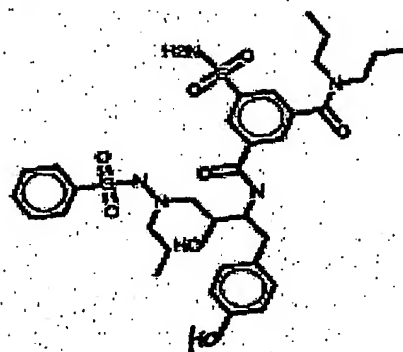
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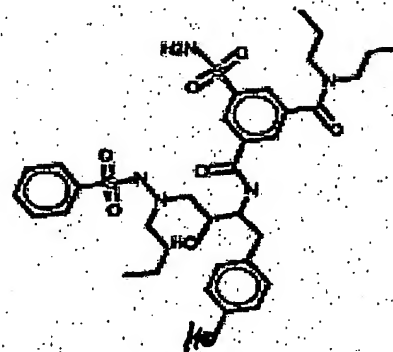
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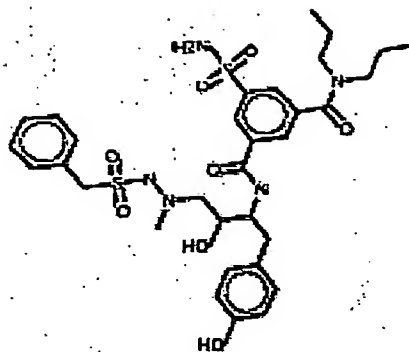
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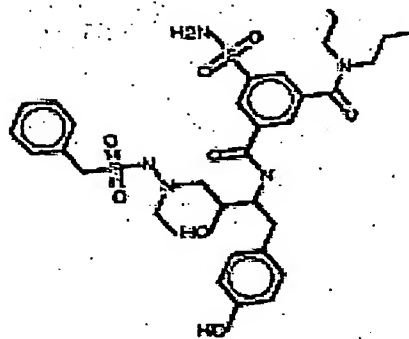
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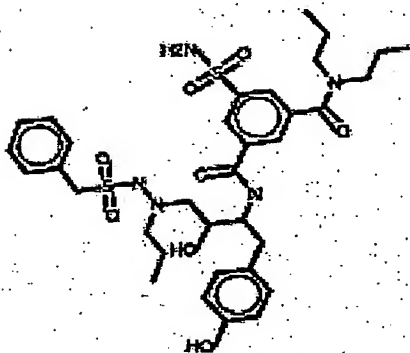
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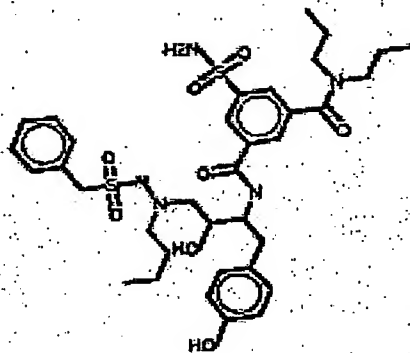
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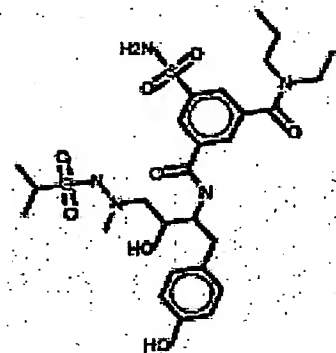
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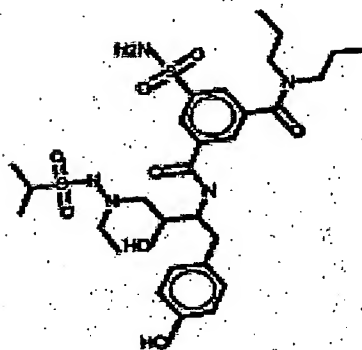
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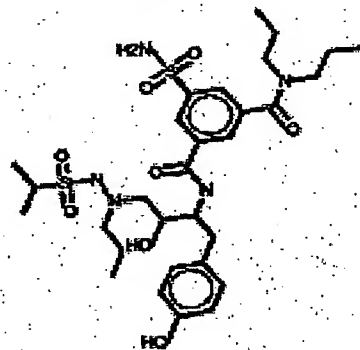
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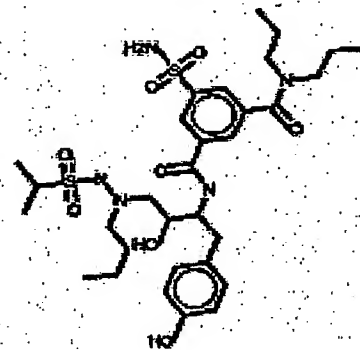
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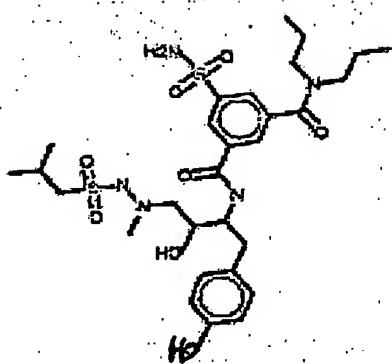
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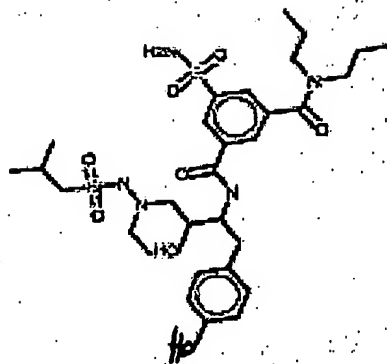
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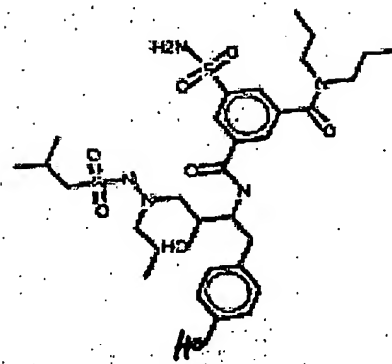
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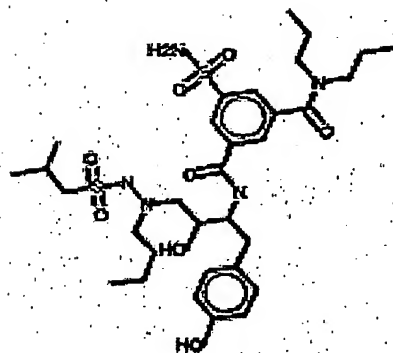
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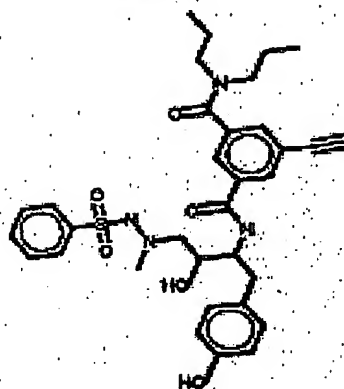
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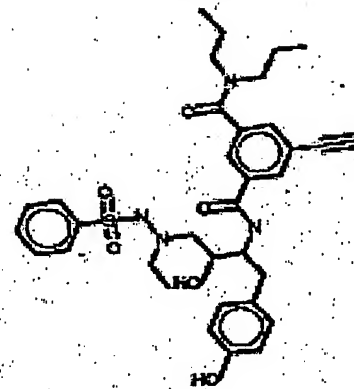
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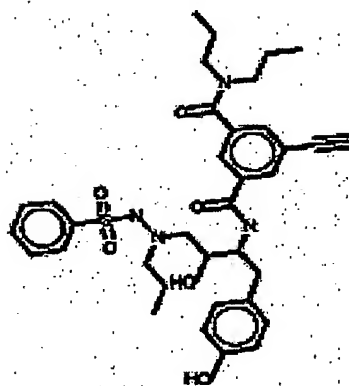
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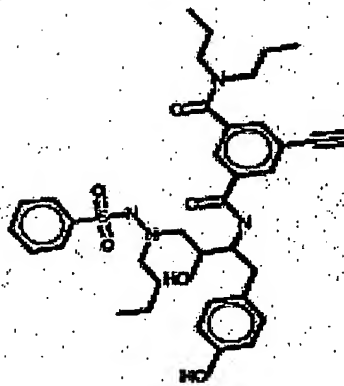
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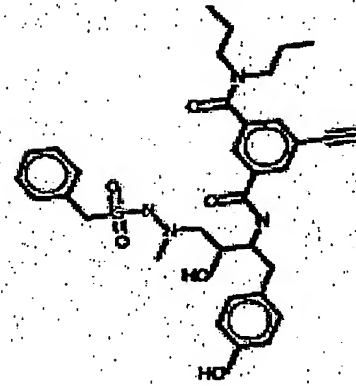
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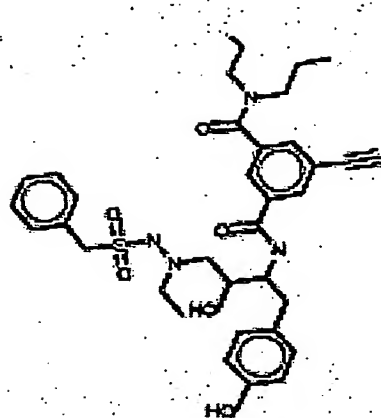
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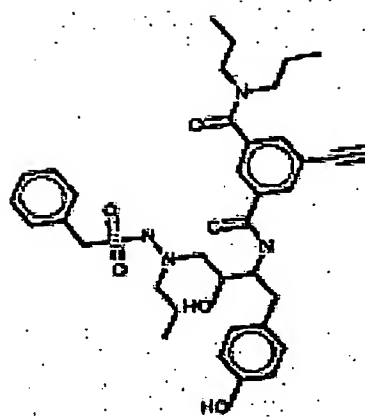
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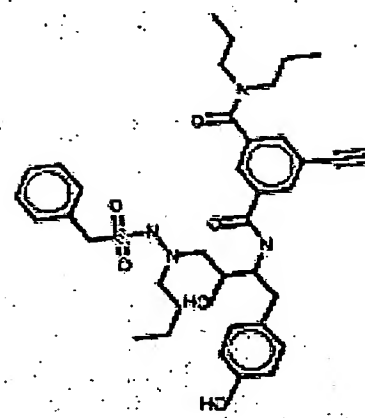
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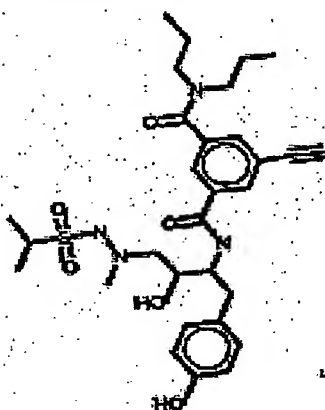
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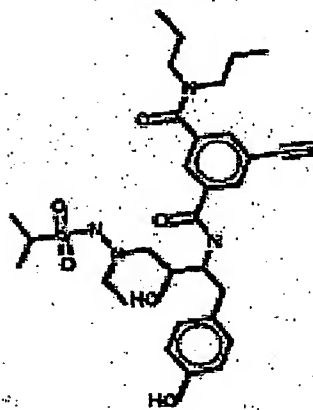
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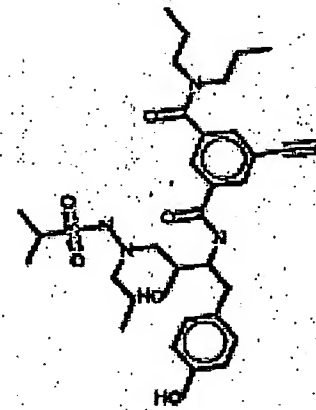
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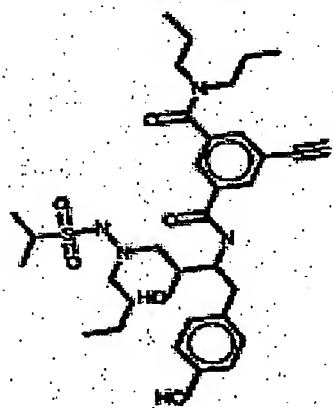
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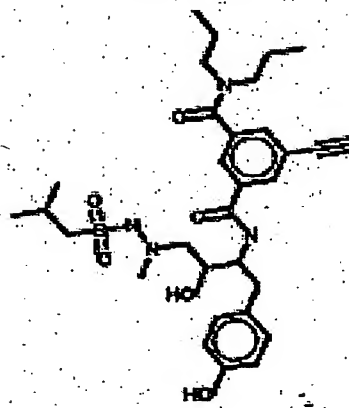
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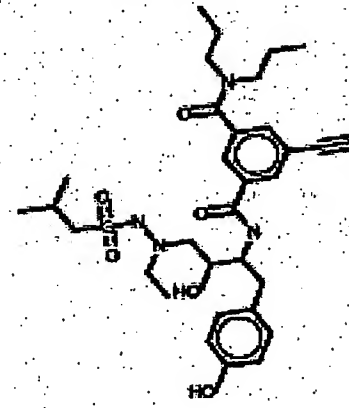
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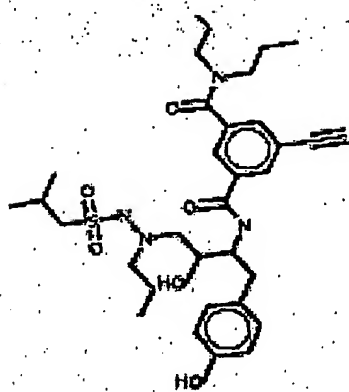
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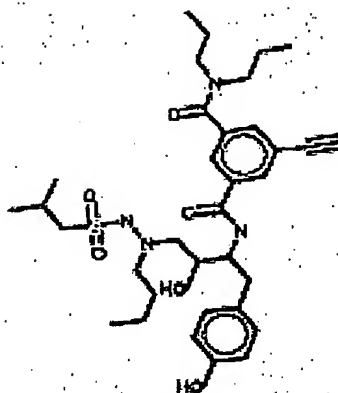
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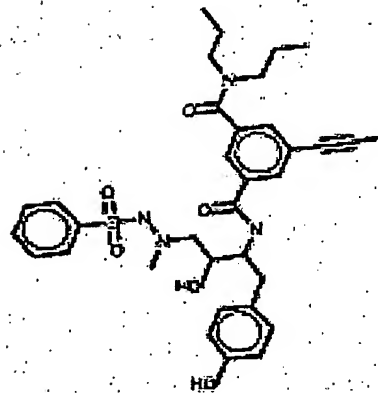
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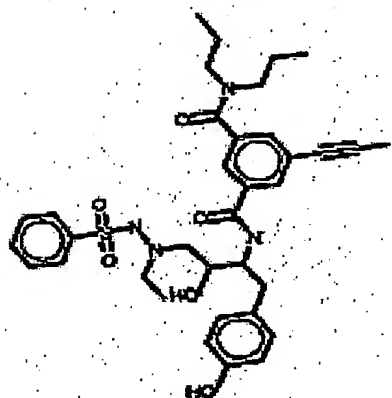
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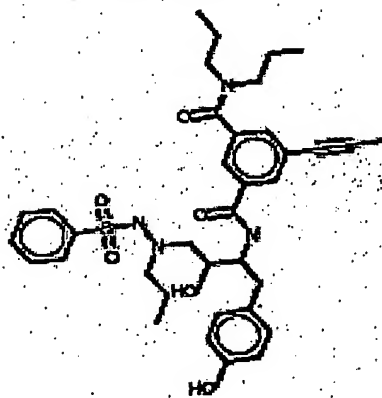
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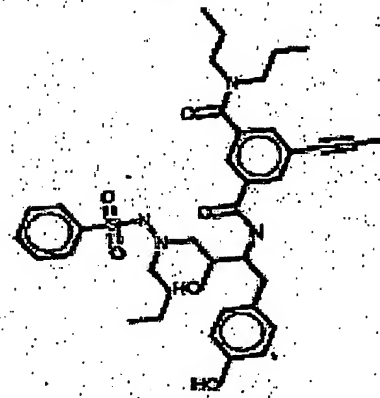
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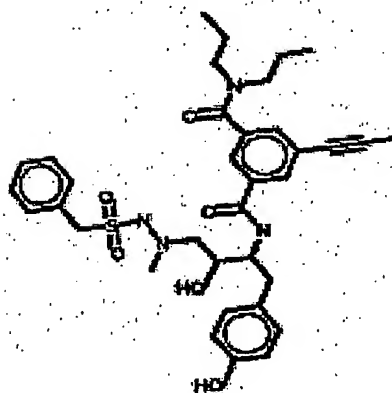
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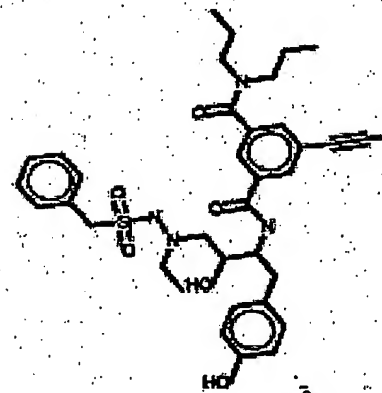
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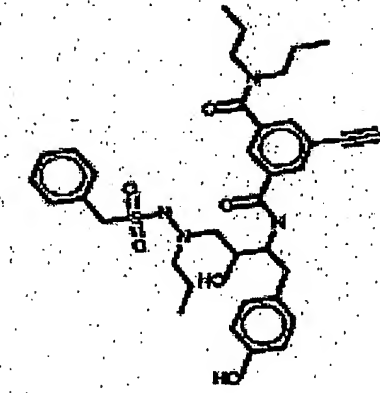
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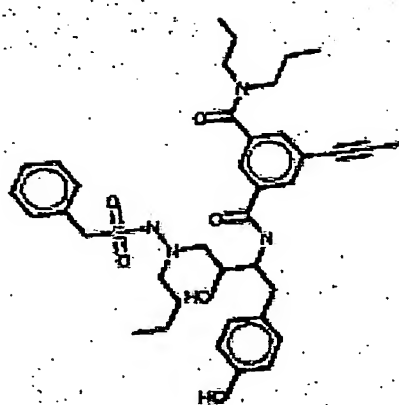
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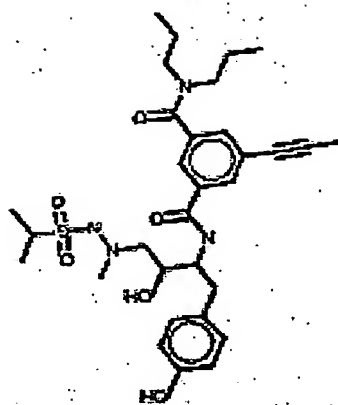
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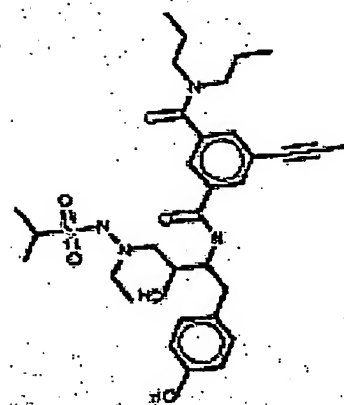
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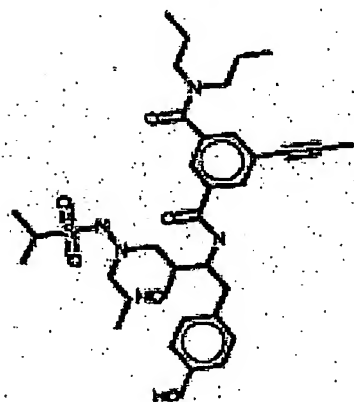
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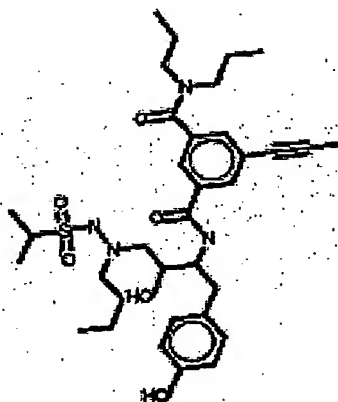
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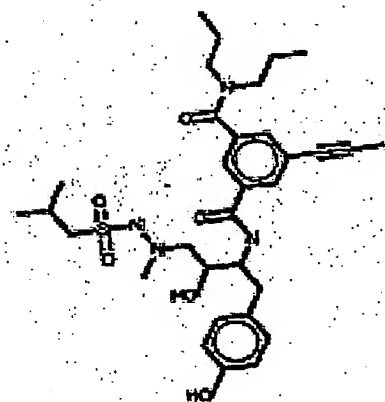
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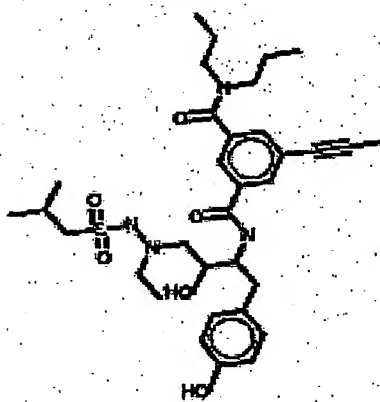
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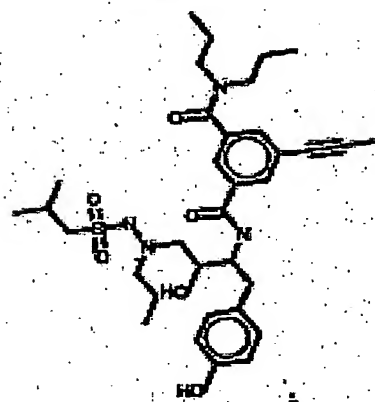
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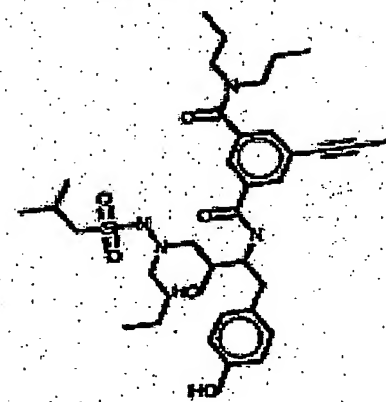
3H9 3.6.2.1



3H10 3.6.2.2



3H11 3.6.2.3



3H12 3.6.2.4

5 The compounds of the invention may exist as geometric or stereoisomers isomers as well as tautomers. Thus, the invention includes all tautomers and geometric isomers, such as the *E* and

Z geometric isomers, as well as mixtures thereof. Furthermore, the invention includes pure enantiomers and diastereomers as well as mixtures thereof, including racemic mixtures. The individual geometric isomers, enantiomers, or diastereomers may
5 be prepared or isolated by methods known in the art.

Compounds of the invention of designated stereochemistry may be included in mixtures, including racemic mixtures, with other enantiomers, diastereomers, geometric isomers or tautomers. Compounds of the invention with designated
10 stereochemistry are typically present in these mixtures in excess of 50 percent. Preferably, compounds of the invention with designated stereochemistry are present in these mixtures in excess of 80 percent. Most preferably, compounds of the invention with designated stereochemistry are present in these
15 mixtures in excess of 90 percent.

Several of the compounds of formula (I) above are amines, and as such form salts when reacted with acids. Pharmaceutically acceptable salts are generally preferred over the corresponding amines of the invention since they typically
20 produce compounds which are more water soluble, stable and/or more crystalline. Pharmaceutically acceptable salts are any salt which retains the activity of the parent compound and does not impart any deleterious or undesirable effect on the subject to whom it is administered and in the context in which it is
25 administered. Pharmaceutically acceptable salts include salts of both inorganic and organic acids. The preferred pharmaceutically acceptable salts include salts of the following acids acetic, aspartic, benzenesulfonic, benzoic, bicarbonic, bisulfuric, bitartaric, butyric, calcium edetate, camsyllic,
30 carbonic, chlorobenzoic, citric, edetic, edisylic, estolic, esyl, esylic, formic, fumaric, gluceptic, gluconic, glutamic, glycollylarsanilic, hexamic, hexylresorcinoic, hydrabamic, hydrobromic, hydrochloric, hydroiodic, hydroxynaphthoic, isethionic, lactic, lactobionic, maleic, malic, malonic,
35 mandelic, methanesulfonic, methylnitric, methylsulfuric, mucic,

muconic, napsylic, nitric, oxalic, p-nitromethanesulfonic, pamoic, pantothenic, phosphoric, monohydrogen phosphoric, dihydrogen phosphoric, phthalic, polygalactouronic, propionic, salicylic, stearic, succinic, succinic, sulfamic, sulfanilic, sulfonic, sulfuric, tannic, tartaric, teoclic and toluenesulfonic. For other acceptable salts, see *Int. J. Pharm.*, 33, 201-217 (1986) and *J. Pharm. Sci.*, 66(1), 1, (1977).

The invention provides compounds, compositions, kits, and methods for inhibiting beta-secretase enzyme activity and A beta peptide production. Inhibition of beta-secretase enzyme activity halts or reduces the production of A beta from APP and reduces or eliminates the formation of beta-amyloid deposits in the brain.

Methods of the Invention

As previously mentioned, compounds of the invention, and pharmaceutically acceptable salts or esters thereof, are useful for treating humans or animals suffering from a condition characterized by a pathological form of beta-amyloid peptide, such as beta-amyloid plaques, and for helping to prevent or delay the onset of such a condition. For example, the compounds are useful for treating Alzheimer's disease, for helping prevent or delay the onset of Alzheimer's disease, for treating patients with MCI (mild cognitive impairment) and preventing or delaying the onset of Alzheimer's disease in those who would progress from MCI to AD, for treating Down's syndrome, for treating humans who have Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, for treating cerebral amyloid angiopathy and preventing its potential consequences, i.e. single and recurrent lobar hemorrhages, for treating other degenerative dementias, including dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's disease, dementia associated with progressive supranuclear palsy, dementia associated with cortical basal degeneration, and diffuse Lewy body type Alzheimer's disease. The compounds and compositions of the invention are particularly useful for treating or

preventing Alzheimer's disease. When treating or preventing these diseases, the compounds of the invention can either be used individually or in combination, as is best for the patient.

To prepare compositions, one or more compounds of the invention are mixed with a suitable pharmaceutically acceptable carrier. Upon mixing or addition of the compound(s), the resulting mixture may be a solution, suspension, emulsion, or the like. Liposomal suspensions may also be suitable as pharmaceutically acceptable carriers. These may be prepared according to methods known to those skilled in the art. The form of the resulting mixture depends upon a number of factors, including the intended mode of administration and the solubility of the compound in the selected carrier or vehicle. The effective concentration is sufficient for lessening or ameliorating at least one symptom of the disease, disorder, or condition treated and may be empirically determined.

Pharmaceutical carriers or vehicles suitable for administration of the compounds provided herein include any such carriers known to those skilled in the art to be suitable for the particular mode of administration. In addition, the active materials can also be mixed with other active materials that do not impair the desired action, or with materials that supplement the desired action, or have another action. The compounds may be formulated as the sole pharmaceutically active ingredient in the composition or may be combined with other active ingredients.

Where the compounds exhibit insufficient solubility, methods for solubilizing may be used. Such methods are known and include, but are not limited to, using cosolvents such as dimethylsulfoxide (DMSO), using surfactants such as Tween®, and dissolution in aqueous sodium bicarbonate. Derivatives of the compounds, such as salts or prodrugs may also be used in formulating effective pharmaceutical compositions.

The concentration of the compound is effective for delivery of an amount upon administration that lessens or ameliorates at

least one symptom of the disorder for which the compound is administered. Typically, the compositions are formulated for single dosage administration.

5 The compounds of the invention may be prepared with carriers that protect them against rapid elimination from the body, such as time-release formulations or coatings. Such carriers include controlled release formulations, such as, but not limited to, microencapsulated delivery systems. The active compound is included in the pharmaceutically acceptable carrier
10 in an amount sufficient to exert a therapeutically useful effect in the absence of undesirable side effects on the patient treated. The therapeutically effective concentration may be determined empirically by testing the compounds in known *in vitro* and *in vivo* model systems for the treated disorder.

15 The compounds and compositions of the invention can be enclosed in multiple or single dose containers. The enclosed compounds and compositions can be provided in kits, for example, including component parts that can be assembled for use. For example, a compound inhibitor in lyophilized form and a suitable
20 diluent may be provided as separated components for combination prior to use. A kit may include a compound inhibitor and a second therapeutic agent for co-administration. The inhibitor and second therapeutic agent may be provided as separate component parts. A kit may include a plurality of containers,
25 each container holding one or more unit dose of the compound of the invention. The containers are preferably adapted for the desired mode of administration, including, but not limited to tablets, gel capsules, sustained-release capsules, and the like for oral administration; depot products, pre-filled syringes,
30 ampoules, vials, and the like for parenteral administration; and patches, medipads, creams, and the like for topical administration.

The concentration of active compound in the drug composition will depend on absorption, inactivation, and
35 excretion rates of the active compound, the dosage schedule, and

amount administered as well as other factors known to those of skill in the art.

The active ingredient may be administered at once, or may be divided into a number of smaller doses to be administered at intervals of time. It is understood that the precise dosage and duration of treatment is a function of the disease being treated and may be determined empirically using known testing protocols or by extrapolation from *in vivo* or *in vitro* test data. It is to be noted that concentrations and dosage values may also vary with the severity of the condition to be alleviated. It is to be further understood that for any particular subject, specific dosage regimens should be adjusted over time according to the individual need and the professional judgment of the person administering or supervising the administration of the compositions, and that the concentration ranges set forth herein are exemplary only and are not intended to limit the scope or practice of the claimed compositions.

If oral administration is desired, the compound should be provided in a composition that protects it from the acidic environment of the stomach. For example, the composition can be formulated in an enteric coating that maintains its integrity in the stomach and releases the active compound in the intestine. The composition may also be formulated in combination with an antacid or other such ingredient.

Oral compositions will generally include an inert diluent or an edible carrier and may be compressed into tablets or enclosed in gelatin capsules. For the purpose of oral therapeutic administration, the active compound or compounds can be incorporated with excipients and used in the form of tablets, capsules, or troches. Pharmaceutically compatible binding agents and adjuvant materials can be included as part of the composition.

The tablets, pills, capsules, troches, and the like can contain any of the following ingredients or compounds of a similar nature: a binder such as, but not limited to, gum

tragacanth, acacia, corn starch, or gelatin; an excipient such as microcrystalline cellulose, starch, or lactose; a disintegrating agent such as, but not limited to, alginic acid and corn starch; a lubricant such as, but not limited to, magnesium stearate; a gildant, such as, but not limited to, colloidal silicon dioxide; a sweetening agent such as sucrose or saccharin; and a flavoring agent such as peppermint, methyl salicylate, or fruit flavoring.

When the dosage unit form is a capsule, it can contain, in addition to material of the above type, a liquid carrier such as a fatty oil. In addition, dosage unit forms can contain various other materials, which modify the physical form of the dosage unit, for example, coatings of sugar and other enteric agents. The compounds can also be administered as a component of an elixir, suspension, syrup, wafer, chewing gum or the like. A syrup may contain, in addition to the active compounds, sucrose as a sweetening agent and certain preservatives, dyes and colorings, and flavors.

The active materials can also be mixed with other active materials that do not impair the desired action, or with materials that supplement the desired action.

Solutions or suspensions used for parenteral, intradermal, subcutaneous, or topical application can include any of the following components: a sterile diluent such as water for injection, saline solution, fixed oil, a naturally occurring vegetable oil such as sesame oil, coconut oil, peanut oil, cottonseed oil, and the like, or a synthetic fatty vehicle such as ethyl oleate, and the like, polyethylene glycol, glycerine, propylene glycol, or other synthetic solvent; antimicrobial agents such as benzyl alcohol and methyl parabens; antioxidants such as ascorbic acid and sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid (EDTA); buffers such as acetates, citrates, and phosphates; and agents for the adjustment of tonicity such as sodium chloride and dextrose. Parenteral preparations can be enclosed in ampoules, disposable

syringes, or multiple dose vials made of glass, plastic, or other suitable material. Buffers, preservatives, antioxidants, and the like can be incorporated as required.

Where administered intravenously, suitable carriers include physiological saline, phosphate buffered saline (PBS), and solutions containing thickening and solubilizing agents such as glucose, polyethylene glycol, polypropyleneglycol, and mixtures thereof. Liposomal suspensions including tissue-targeted liposomes may also be suitable as pharmaceutically acceptable carriers. These may be prepared according to methods known for example, as described in U.S. Patent No. 4,522,811.

The active compounds may be prepared with carriers that protect the compound against rapid elimination from the body, such as time-release formulations or coatings. Such carriers include controlled release formulations, such as, but not limited to, implants and microencapsulated delivery systems, and biodegradable, biocompatible polymers such as collagen, ethylene vinyl acetate, polyanhydrides, polyglycolic acid, polyorthoesters, polylactic acid, and the like. Methods for preparation of such formulations are known to those skilled in the art.

The compounds of the invention can be administered orally, parenterally (IV, IM, depo-IM, SQ, and depo-SQ), sublingually, intranasally (inhalation), intrathecally, topically, or rectally. Dosage forms known to those skilled in the art are suitable for delivery of the compounds of the invention.

Compounds of the invention may be administered enterally or parenterally. When administered orally, compounds of the invention can be administered in usual dosage forms for oral administration as is well known to those skilled in the art. These dosage forms include the usual solid unit dosage forms of tablets and capsules as well as liquid dosage forms such as solutions, suspensions, and elixirs. When the solid dosage forms are used, it is preferred that they be of the sustained

release type so that the compounds of the invention need to be administered only once or twice daily.

The oral dosage forms are administered to the patient 1, 2, 3, or 4 times daily. It is preferred that the compounds of the invention be administered either three or fewer times, more preferably once or twice daily. Hence, it is preferred that the compounds of the invention be administered in oral dosage form. It is preferred that whatever oral dosage form is used, that it be designed so as to protect the compounds of the invention from the acidic environment of the stomach. Enteric coated tablets are well known to those skilled in the art. In addition, capsules filled with small spheres each coated to protect from the acidic stomach, are also well known to those skilled in the art.

When administered orally, an administered amount therapeutically effective to inhibit beta-secretase activity, to inhibit A beta production, to inhibit A beta deposition, or to treat or prevent AD is from about 0.1 mg/day to about 1,000 mg/day. It is preferred that the oral dosage is from about 1 mg/day to about 100 mg/day. It is more preferred that the oral dosage is from about 5 mg/day to about 50 mg/day. It is understood that while a patient may be started at one dose, that dose may be varied over time as the patient's condition changes.

Compounds of the invention may also be advantageously delivered in a nano crystal dispersion formulation. Preparation of such formulations is described, for example, in U.S. Patent 5,145,684. Nano crystalline dispersions of HIV protease inhibitors and their method of use are described in U.S. Patent No. 6,045,829. The nano crystalline formulations typically afford greater bioavailability of drug compounds.

The compounds of the invention can be used in combination, with each other or with other therapeutic agents or approaches used to treat or prevent the conditions listed above. Such agents or approaches include: acetylcholine esterase inhibitors such as tacrine (tetrahydroaminoacridine, marketed as COGNEX®),

donepezil hydrochloride, (marketed as Aricept® and rivastigmine (marketed as Exelon®); gamma-secretase inhibitors; anti-inflammatory agents such as cyclooxygenase II inhibitors; anti-oxidants such as Vitamin E and ginkgolides; immunological approaches, such as, for example, immunization with A beta peptide or administration of anti-A beta peptide antibodies; statins; and direct or indirect neurotropic agents such as Cerebrolysin®, AIT-082 (Emilieu, 2000, Arch. Neurol. 57:454), and other neurotropic agents of the future.

10 It should be apparent to one skilled in the art that the exact dosage and frequency of administration will depend on the particular compounds of the invention administered, the particular condition being treated, the severity of the condition being treated, the age, weight, general physical
15 condition of the particular patient, and other medication the individual may be taking as is well known to administering physicians who are skilled in this art.

The invention may be further understood with reference to the following biological examples. These examples are intended
20 to be representative of specific embodiments of the invention, and are not intended as limiting the scope of the invention.

BIOLOGY EXAMPLES

Biology Example A

Enzyme Inhibition Assay

25 The compounds of the invention are analyzed for inhibitory activity by use of the MBP-C125 assay. This assay determines the relative inhibition of beta-secretase cleavage of a model APP substrate, MBP-C125SW, by the compounds assayed as compared with an untreated control. A detailed description of the assay
30 parameters can be found, for example, in U.S. Patent No. 5,942,400. Briefly, the substrate is a fusion peptide formed of maltose binding protein (MBP) and the carboxy terminal 125 amino acids of APP-SW, the Swedish mutation. The beta-secretase enzyme is derived from human brain tissue as described in Sinha
35 et al, 1999, Nature 40:537-540) or recombinantly produced as the

full-length enzyme (amino acids 1-501), and can be prepared, for example, from 293 cells expressing the recombinant cDNA, as described in WO00/47618.

Inhibition of the enzyme is analyzed, for example, by immunoassay of the enzyme's cleavage products. One exemplary ELISA uses an anti-MBP capture antibody that is deposited on precoated and blocked 96-well high binding plates, followed by incubation with diluted enzyme reaction supernatant, incubation with a specific reporter antibody, for example, biotinylated anti-SW192 reporter antibody, and further incubation with streptavidin/alkaline phosphatase. In the assay, cleavage of the intact MBP-C125SW fusion protein results in the generation of a truncated amino-terminal fragment, exposing a new SW-192 antibody-positive epitope at the carboxy terminus. Detection is effected by a fluorescent substrate signal on cleavage by the phosphatase. ELISA only detects cleavage following Leu 596 at the substrate's APP-SW 751 mutation site.

Specific Assay Procedure:

Compounds are diluted in a 1:1 dilution series to a six-point concentration curve (two wells per concentration) in one 96-plate row per compound tested. Each of the test compounds is prepared in DMSO to make up a 10 millimolar stock solution. The stock solution is serially diluted in DMSO to obtain a final compound concentration of 200 micromolar at the high point of a 6-point dilution curve. Ten (10) microliters of each dilution is added to each of two wells on row C of a corresponding V-bottom plate to which 190 microliters of 52 millimolar NaOAc, 7.9% DMSO, pH 4.5 are pre-added. The NaOAc diluted compound plate is spun down to pellet precipitant and 20 microliters/well is transferred to a corresponding flat-bottom plate to which 30 microliters of ice-cold enzyme-substrate mixture (2.5 microliters MBP-C125SW substrate, 0.03 microliters enzyme and 24.5 microliters ice cold 0.09% TX100 per 30 microliters) is added. The final reaction mixture of 200 micromolar compound at

the highest curve point is in 5% DMSO, 20 millimolar NaOAc, 0.06% TX100, at pH 4.5.

Warming the plates to 37 degrees C starts the enzyme reaction. After 90 minutes at 37 degrees C, 200
5 microliters/well cold specimen diluent is added to stop the reaction and 20 microliters/well was transferred to a corresponding anti-MBP antibody coated ELISA plate for capture, containing 80 microliters/well specimen diluent. This reaction is incubated overnight at 4 degrees C and the ELISA is developed
10 the next day after a 2 hour incubation with anti-192SW antibody, followed by Streptavidin-AP conjugate and fluorescent substrate. The signal is read on a fluorescent plate reader.

Relative compound inhibition potency is determined by calculating the concentration of compound that showed a fifty
15 percent reduction in detected signal (IC_{50}) compared to the enzyme reaction signal in the control wells with no added compound. In this assay, the compounds of the invention exhibited an IC_{50} of less than 50 micromolar.

Biology Example B

20 Cell Free Inhibition Assay Utilizing a Synthetic APP Substrate

A synthetic APP substrate that can be cleaved by beta-secretase and having N-terminal biotin and made fluorescent by the covalent attachment of Oregon green at the Cys residue is used to assay beta-secretase activity in the presence or absence
25 of the inhibitory compounds of the invention. Useful substrates include the following:

Biotin-SEVNL-DAEFR [Oregon green] KK	[SEQ ID NO: 1]
Biotin-SEVKM-DAEFR [Oregon green] KK	[SEQ ID NO: 2]
30 Biotin-GLNIKTEEISEISY-EVEFRC [Oregon green] KK	[SEQ ID NO: 3]
Biotin-ADRGLTTRPGSGLTNIKTEEISEVNL-DAEF [Oregon green] KK	[SEQ ID NO: 4]
Biotin-FVNQHLCoxGSHLVEALY-LVCoxGERGFFYTPKA [Oregon green] KK	[SEQ ID NO: 5]

The enzyme (0.1 nanomolar) and test compounds (0.001 - 100 micromolar) are incubated in pre-blocked, low affinity, black plates (384 well) at 37 degrees for 30 minutes. The reaction is initiated by addition of 150 millimolar substrate to a final volume of 30 microliter per well. The final assay conditions are: 0.001 - 100 micromolar compound inhibitor; 0.1 molar sodium acetate (pH 4.5); 150 nanomolar substrate; 0.1 nanomolar soluble beta-secretase; 0.001% Tween 20, and 2% DMSO. The assay mixture is incubated for 3 hours at 37 degrees C, and the reaction is terminated by the addition of a saturating concentration of immunopure streptavidin. After incubation with streptavidin at room temperature for 15 minutes, fluorescence polarization is measured, for example, using a LJL Acquest (Ex485 nm/ Em530 nm). The activity of the beta-secretase enzyme is detected by changes in the fluorescence polarization that occur when the substrate is cleaved by the enzyme. Incubation in the presence or absence of compound inhibitor demonstrates specific inhibition of beta-secretase enzymatic cleavage of its synthetic APP substrate. In this assay, compounds of the invention exhibited an IC₅₀ of less than 50 micromolar.

Biology Example C

Beta-Secretase Inhibition: P26-P4'SW Assay

Synthetic substrates containing the beta-secretase cleavage site of APP are used to assay beta-secretase activity, using the methods described, for example, in published PCT application WO00/47618. The P26-P4'SW substrate is a peptide of the sequence:

(biotin) CGGADRGLTTRPGSGLTNIKTEEISEVNLD AEF [SEQ ID NO: 6]

The P26-P1 standard has the sequence:

(biotin) CGGADRGLTTRPGSGLTNIKTEEISEVNL [SEQ ID NO: 7].

Briefly, the biotin-coupled synthetic substrates are incubated at a concentration of from about 0 to about 200 micromolar in this assay. When testing inhibitory compounds, a substrate concentration of about 1.0 micromolar is preferred.

5 Test compounds diluted in DMSO are added to the reaction mixture, with a final DMSO concentration of 5%. Controls also contain a final DMSO concentration of 5%. The concentration of beta secretase enzyme in the reaction is varied, to give product concentrations with the linear range of the ELISA assay, about

10 125 to 2000 picomolar, after dilution.

The reaction mixture also includes 20 millimolar sodium acetate, pH 4.5, 0.06% Triton X100, and is incubated at 37 degrees C for about 1 to 3 hours. Samples are then diluted in assay buffer (for example, 145.4 nanomolar sodium chloride, 9.51

15 millimolar sodium phosphate, 7.7 millimolar sodium azide, 0.05% Triton X405, 6g/liter bovine serum albumin, pH 7.4) to quench the reaction, then diluted further for immunoassay of the cleavage products.

Cleavage products can be assayed by ELISA. Diluted samples

20 and standards are incubated in assay plates coated with capture antibody, for example, SW192, for about 24 hours at 4 degrees C. After washing in TTBS buffer (150 millimolar sodium chloride, 25 millimolar Tris, 0.05% Tween 20, pH 7.5), the samples are incubated with streptavidin-AP according to the manufacturer's

25 instructions. After a one hour incubation at room temperature, the samples are washed in TTBS and incubated with fluorescent substrate solution A (31.2 g/liter 2-amino-2-methyl-1-propanol, 30 mg/liter, pH 9.5). Reaction with streptavidin-alkaline phosphate permits detection by fluorescence. Compounds that are

30 effective inhibitors of beta-secretase activity demonstrate reduced cleavage of the substrate as compared to a control.

Biology Example D

Assays using Synthetic Oligopeptide-Substrates

Synthetic oligopeptides are prepared that incorporate the

35 known cleavage site of beta-secretase, and optionally detectable

tags, such as fluorescent or chromogenic moieties. Examples of such peptides, as well as their production and detection methods are described in U.S. Patent No: 5,942,400, herein incorporated by reference. Cleavage products can be detected using high performance liquid chromatography, or fluorescent or chromogenic detection methods appropriate to the peptide to be detected, according to methods well known in the art.

By way of example, one such peptide has the sequence SEVNLDAEF [SEQ ID NO: 8], and the cleavage site is between residues 5 and 6. Another preferred substrate has the sequence ADRLTTRPGSGLTNIKTEEISEVNL-DAEF [SEQ ID NO: 9], and the cleavage site is between residues 26 and 27.

These synthetic APP substrates are incubated in the presence of beta-secretase under conditions sufficient to result in beta-secretase mediated cleavage of the substrate. Comparison of the cleavage results in the presence of the compound inhibitor to control results provides a measure of the compound's inhibitory activity.

Biology Example E

20 Inhibition of Beta-Secretase Activity - Cellular Assay

An exemplary assay for the analysis of inhibition of beta-secretase activity utilizes the human embryonic kidney cell line HEKp293 (ATCC Accession No. CRL-1573) transfected with APP751 containing the naturally occurring double mutation Lys651Met52 to Asn651Leu652 (numbered for APP751), commonly called the Swedish mutation and shown to overproduce A beta (Citron et al., 1992, Nature 360:672-674), as described in U.S. Patent No. 5,604,102.

The cells are incubated in the presence/absence of the inhibitory compound (diluted in DMSO) at the desired concentration, generally up to 10 micrograms/ml. At the end of the treatment period, conditioned media is analyzed for beta-secretase activity, for example, by analysis of cleavage fragments. A beta can be analyzed by immunoassay, using specific detection antibodies. The enzymatic activity is

measured in the presence and absence of the compound inhibitors to demonstrate specific inhibition of beta-secretase mediated cleavage of APP substrate.

Biology Example F

5 Inhibition of Beta-Secretase in Animal Models of AD

Various animal models can be used to screen for inhibition of beta-secretase activity. Examples of animal models useful in the invention include, but are not limited to, mouse, guinea pig, dog, and the like. The animals used can be wild type, 10 transgenic, or knockout models. In addition, mammalian models can express mutations in APP, such as APP695-SW and the like described herein. Examples of transgenic non-human mammalian models are described in U.S. Patent Nos. 5,604,102, 5,912,410 and 5,811,633.

15 PDAPP mice, prepared as described in Games et al., 1995, Nature 373:523-527 are useful to analyze *in vivo* suppression of A beta release in the presence of putative inhibitory compounds. As described in U.S. Patent No. 6,191,166, 4 month old PDAPP mice are administered compound formulated in vehicle, such as 20 corn oil. The mice are dosed with compound (1-30 mg/ml; preferably 1-10 mg/ml). After time, e.g., 3-10 hours, the animals are sacrificed, and brains removed for analysis.

Transgenic animals are administered an amount of the compound inhibitor formulated in a carrier suitable for the 25 chosen mode of administration. Control animals are untreated, treated with vehicle, or treated with an inactive compound. Administration can be acute, i.e., single dose or multiple doses in one day, or can be chronic, i.e., dosing is repeated daily for a period of days. Beginning at time 0, brain tissue or 30 cerebral fluid is obtained from selected animals and analyzed for the presence of APP cleavage peptides, including A beta, for example, by immunoassay using specific antibodies for A beta detection. At the end of the test period, animals are sacrificed and brain tissue or cerebral fluid is analyzed for

the presence of A beta and/or beta-amyloid plaques. The tissue is also analyzed for necrosis.

Animals administered the compound inhibitors of the invention are expected to demonstrate reduced A beta in brain tissues or cerebral fluids and reduced beta amyloid plaques in brain tissue, as compared with non-treated controls.

Biology Example G

Inhibition of A Beta Production in Human Patients

Patients suffering from Alzheimer's Disease (AD) demonstrate an increased amount of A beta in the brain. AD patients are administered an amount of the compound inhibitor formulated in a carrier suitable for the chosen mode of administration. Administration is repeated daily for the duration of the test period. Beginning on day 0, cognitive and memory tests are performed, for example, once per month.

Patients administered the compound inhibitors are expected to demonstrate slowing or stabilization of disease progression as analyzed by changes in one or more of the following disease parameters: A beta present in CSF or plasma; brain or hippocampal volume; A beta deposits in the brain; amyloid plaque in the brain; and scores for cognitive and memory function, as compared with control, non-treated patients.

Biology Example H

Prevention of A Beta Production in Patients at Risk for AD

Patients predisposed or at risk for developing AD are identified either by recognition of a familial inheritance pattern, for example, presence of the Swedish Mutation, and/or by monitoring diagnostic parameters. Patients identified as predisposed or at risk for developing AD are administered an amount of the compound inhibitor formulated in a carrier suitable for the chosen mode of administration. Administration is repeated daily for the duration of the test period. Beginning on day 0, cognitive and memory tests are performed, for example, once per month.

Patients administered the compound inhibitors are expected to demonstrate slowing or stabilization of disease progression as analyzed by changes in one or more of the following disease parameters: A beta present in CSF or plasma; brain or hippocampal volume; amyloid plaque in the brain; and scores for cognitive and memory function, as compared with control, non-treated patients.

It should be noted that, as used in this specification and the appended claims, the singular forms "a," "an," and "the" include plural referents unless the content clearly dictates otherwise. Thus, for example, reference to a composition containing "a compound" includes a mixture of two or more compounds. It should also be noted that the term "or" is generally employed in its sense including "and/or" unless the content clearly dictates otherwise.

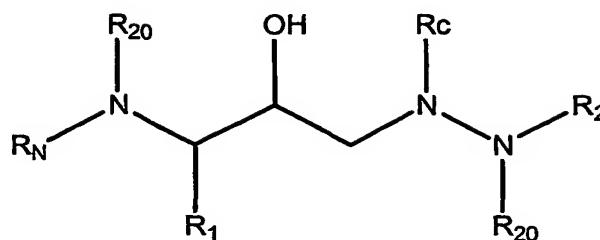
Unless defined otherwise, all scientific and technical terms used herein have the same meaning as commonly understood by one of skill in the art to which this invention belongs.

All patents and publications referred to herein are hereby incorporated by reference for all purposes.

The invention has been described with reference to various specific and preferred embodiments and techniques. However, it should be understood that many variations and modifications may be made while remaining within the spirit and scope of the invention.

WHAT IS CLAIMED IS:

1. A compound of the formula II:



(II)

or a pharmaceutically acceptable salt thereof,
where Rc is

- (I) -C₁-C₁₀ alkyl optionally substituted with one, two or three groups independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, -NR_{1-a}R_{1-b}, -OC(=O)NR_{1-a}R_{1-b}, -S(=O)₀₋₂R_{1-a}, -NR_{1-a}C(=O)NR_{1-a}R_{1-b}, -C(=O)NR_{1-a}R_{1-b}, and -S(=O)₂NR_{1-a}R_{1-b} wherein

- R_{1-a} and R_{1-b} at each occurrence are independently H or C₁-C₆ alkyl,

- (II) -(CH₂)₀₋₃-(C₃-C₈) cycloalkyl where cycloalkyl can be optionally substituted with one, two or three substituents independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, -CO₂H, -CO₂-(C₁-C₄ alkyl), and -NR_{1-a}R_{1-b}

- (III) -(CR_{C-x}R_{C-y})₀₋₄-R_{C-aryl} where R_{C-x} and R_{C-y} are independently selected from the group consisting of

- H,
C₁-C₄ alkyl optionally substituted with 1 or 2 -OH,
C₁-C₄ alkoxy optionally substituted with 1, 2, or 3 halogen,
-(CH₂)₀₋₄-C₃-C₈ cycloalkyl,
C₂-C₆ alkenyl containing one or two double bonds,
C₂-C₆ alkynyl containing one or two triple bonds, and
phenyl,

or

R_{C-x} and R_{C-y} are taken together with the carbon to which they are attached to form a carbocycle of three, four, five, six or seven carbon atoms, where one carbon atom is optionally replaced by a group selected from -O-, -S-, -SO₂-, -NR_{N-2}- and R_{C-aryl} , wherein

R_{C-aryl} is phenyl, which is optionally substituted with 1, 2, or 3 groups that are independently:

(1) C₁-C₆ alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(2) -OH,

(3) -NO₂,

(4) halogen,

(5) -CO₂H,

(6) -C≡N,

(7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3} where

R_{N-2} and R_{N-3} are independently selected from the group consisting of:

(a) -H,

(b) -C₁-C₆ alkyl optionally substituted with one substituent selected from the group consisting of:

(i) -OH, and

(ii) -NH₂,

(c) -C₁-C₆ alkyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -Br, -I, or OH,

(d) -C₃-C₇ cycloalkyl,

(e) -(C₁-C₂ alkyl)-(C₃-C₇ cycloalkyl),

(f) -(C₁-C₆ alkyl)-O-(C₁-C₃ alkyl),

(g) -C₂-C₆ alkenyl

(h) -C₂-C₆ alkynyl

(i) -C₁-C₆ alkyl chain with one double bond and one triple bond,

(j) $-R_{1-aryl}$ wherein R_{1-aryl} at each occurrence is independently phenyl, naphthyl, indanyl, indenyl, dihydronaphthyl, or tetralinyl each of which is optionally substituted with 1, 2, 3, or 4 groups that are independently:

5 (i) C_1-C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

10 (ii) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

15 (iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(iv) $-F$, Cl , $-Br$ and $-I$,

(v) $-C_1-C_6$ alkoxy optionally substituted with 1, 2, or 3 $-F$,

20 (vi) $-NR_{N-2}R_{N-3}$,

(vii) $-OH$,

(viii) $-C\equiv N$,

(ix) C_3-C_7 cycloalkyl, optionally substituted with 1, 2, or 3 groups that are selected from the
25 group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(x) $-CO-(C_1-C_4 \text{ alkyl})$,

(xi) $-SO_2-NR_{1-a}R_{1-b}$,

(xii) $-CO-NR_{1-a}R_{1-b}$, or

30 (xiii) $-SO_2-(C_1-C_4 \text{ alkyl})$,

(k) $-R_{1-heteroaryl}$ wherein $R_{1-heteroaryl}$ at each occurrence is independently selected from the group consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl,

isoquinolyl, quinazoliny, quinoxaliny, phthalaziny,
imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl,
indoliziny, indazolyl, benzothiazolyl, benzimidazolyl,
benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl,
5 thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridiny,
imidazopyridiny, isothiazolyl, naphthyridiny, cinnoliny,
carbazolyl, beta-carboliny, isochromanyl, chromanyl,
tetrahydroisoquinoliny, isoindoliny,
isobenzotetrahydrofuranyl, isobenzotetrahydrothienyl,
10 isobenzothienyl, benzoxazolyl, pyridopyridiny,
benzotetrahydrofuranyl, benzotetrahydrothienyl, puriny,
benzodioxolyl, triaziny, phenoxaziny, phenothiaziny,
pteridiny, benzothiazolyl, imidazopyridiny, imidazothiazolyl,
dihydrobenzisoxaziny, benzisoxaziny, benzoxaziny,
15 dihydrobenzisothiaziny, benzopyranyl, benzothiopyranyl,
coumariny, isocoumariny, chromony, chromanony, pyridiny-N-
oxide, tetrahydroquinoliny, dihydroquinoliny,
dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumariny,
dihydroisocoumariny, isoindolinonyl, benzodioxanyl,
20 benzoxazolinonyl, pyrrolyl N-oxide, pyrimidiny N-oxide,
pyridaziny N-oxide, pyraziny N-oxide, quinoliny N-oxide,
indolyl N-oxide, indoliny N-oxide, isoquinolyl N-oxide,
quinazoliny N-oxide, quinoxaliny N-oxide, phthalaziny N-
oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
25 thiazolyl N-oxide, indoliziny N-oxide, indazolyl N-oxide,
benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and
benzothiopyranyl S,S-dioxide,

30 where the R₁-heteroaryl group is
optionally substituted with 1, 2, 3, or 4 groups that are
independently:

(i) C₁-C₆ alkyl optionally
substituted with 1, 2, or 3 groups independently selected from

the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C₁-C₃ alkoxy,

(ii) C₂-C₆ alkenyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(iii) C₂-C₆ alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(iv) -F, -Cl, -Br and -I,

(v) -C₁-C₆ alkoxy optionally substituted with one, two, or three -F,

(vi) -(CH₂)₀₋₄-NR_{N-2}R_{N-3},

(vii) -OH,

(viii) -C≡N,

(ix) (CH₂)₀₋₄-C₃-C₇ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(x) (CH₂)₀₋₄-CO-(C₁-C₆ alkyl),

(xi) (CH₂)₀₋₄-SO₂-NR_{N-2}R_{N-3},

(xii) (CH₂)₀₋₄-CO-NR_{N-2}R_{N-3},

(xiii) (CH₂)₀₋₄-SO₂-(C₁-C₆

alkyl),

(xiv) (CH₂)₀₋₄-N(R_{N-2})-SO₂-,

and

(xv) (CH₂)₀₋₄-N(R_{N-2})-C(O)-,

(8) -(CH₂)₀₋₄-CO-(C₁-C₁₂ alkyl),

(9) -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkenyl),

(10) -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkynyl),

(11) -(CH₂)₀₋₄-CO-(CH₂)₀₋₄ (C₃-C₇ cycloalkyl),

(12) -(CH₂)₀₋₄-CO-R₁-aryl,

(13) -(CH₂)₀₋₄-CO-R₁-heteroaryl,

(14) -(CH₂)₀₋₄-CO-R₁-heterocycle wherein

R_1 -heterocycle at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, 5 tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl, dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl, 10 dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the R_1 -heterocycle group is bonded by any atom of the parent R_1 -heterocycle group substituted by hydrogen such that the new bond to the R_1 -heterocycle group replaces the 15 hydrogen atom and its bond, where heterocycle is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(a) C_1 - C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1 - C_3 alkyl, halogen, -OH, 20 -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1 - C_3 alkoxy,

(b) C_2 - C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$ 25

(c) C_2 - C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1 - C_3 alkoxy, and $-NR_{1-a}R_{1-b}$ 30

(d) halogen,

(e) C_1 - C_6 alkoxy,

(f) $-C_1$ - C_6 alkoxy optionally substituted with one, two, or three -F,

(g) $-NR_{N-2}R_{N-3}$,

(h) -OH,

(i) $-C\equiv N$,
 (j) $(CH_2)_{0-4}-(C_3-C_7 \text{ cycloalkyl})$,
 optionally substituted with 1, 2, or 3 groups independently
 selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-$
 5 CF_3 , C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(k) $-(CH_2)_{0-4}-CO-(C_1-C_4 \text{ alkyl})$,

(l) $-(CH_2)_{0-4}-SO_2-NR_{1-a}R_{1-b}$,

(m) $-(CH_2)_{0-4}-CO-NR_{1-a}R_{1-b}$,

(n) $-(CH_2)_{0-4}-SO_2-(C_1-C_6 \text{ alkyl})$, and

10

(o) $=O$,

(p) $-(CH_2)_{0-4}-N(R_{N-2})-SO_2-$

(q) $-(CH_2)_{0-4}-N(R_{N-2})-C(O)-$

(15) $-(CH_2)_{0-4}-CO-R_{N-4}$ wherein

15 R_{N-4} at each occurrence is independently
 selected from the group consisting of morpholinyl,
 thiomorpholinyl, pyrrolidinonyl, pyrrolyl, pyrazolyl, thienyl,
 pyridyl N-oxide, piperazinyl, piperidinyl, homomorpholinyl,
 homothiomorpholinyl, homothiomorpholinyl S-oxide,
 homothiomorpholinyl S,S-dioxide, pyrrolinyl and pyrrolidinyl
 20 where each group is optionally substituted with 1, 2, 3, or 4
 groups that are independently C_1-C_6 alkyl,

(16) $-(CH_2)_{0-4}-CO_2-R_{N-5}$ where

R_{N-5} at each occurrence is independently
 selected from the group consisting of:

25

(a) C_1-C_6 alkyl,

(b) $-(CH_2)_{0-2}-(R_{1-aryl})$,

(c) C_2-C_6 alkenyl,

(d) C_2-C_6 alkynyl,

(e) C_3-C_7 cycloalkyl, and

30

(f) $-(CH_2)_{0-4}-(R_{1-heteroaryl})$,

(17) $-(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$

(18) $-(CH_2)_{0-4}-SO-(C_1-C_8 \text{ alkyl})$,

(19) $-(CH_2)_{0-4}-SO_2-(C_1-C_{12} \text{ alkyl})$,

(20) $-(CH_2)_{0-4}-SO_2-(C_3-C_7 \text{ cycloalkyl})$,

35

(21) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO_2-R_{N-5}$,

- (22) - $(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (23) - $(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (24) - $(\text{CH}_2)_{0-4}-\text{N}(-\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2}$,
 (25) - $(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
 5 (26) - $(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4}$,
 (27) - $(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
 (28) - $(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$ where R_{100} is
 independently H or C_1-C_4 alkyl,
 (29) - $(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
 10 (30) - $(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (31) - $(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
 (32) - $(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
 (33) - $(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
 (34) - $(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl})$ wherein the
 15 alkyl group is optionally substituted with one, two, three,
 four, or five substituents independently selected from the group
 consisting of F, Cl, Br, and I,
 (35) - $(\text{CH}_2)_{0-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
 (36) C_2-C_6 alkenyl optionally substituted
 20 with C_1-C_3 alkyl, halogen, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, or
 $-\text{NR}_{1-a}\text{R}_{1-b}$,
 (37) C_2-C_6 alkynyl optionally substituted
 with C_1-C_3 alkyl, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3
 alkoxy, or $-\text{NR}_{1-a}\text{R}_{1-b}$, and
 25 (38) - $(\text{CH}_2)_{0-4}-\text{N}(-\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$;
 (IV) - $(\text{CR}_{\text{C}-x}\text{R}_{\text{C}-y})_{0-4}-\text{R}_{\text{C-heteroaryl}}$ wherein $\text{R}_{\text{C-heteroaryl}}$ at each
 occurrence is independently selected from the group consisting
 of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl,
 indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl,
 30 quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl,
 pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl,
 benzoisothiazolyl, benzimidazolyl, benzofuranyl, furanyl,
 thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl,
 tetrazolyl, oxazolopyridinyl, isothiazolyl, naphthyridinyl,
 35 cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl,

tetrahydroisoquinolinyl, isoindolinyl,
 isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl,
 isobenzothienyl, benzoxazolyl, pyridopyridinyl,
 benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl,
 5 benzodioxolyl, triazinyl, henoxazinyl, phenothiazinyl,
 pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
 dihydrobenzisoaxazinyl, benzisoxazinyl, benzoxazinyl,
 dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl,
 coumarinyl, isocoumarinyl, chromonyl, chromanonyl,
 10 tetrahydroquinolinyl, dihydroquinolinyl,
 dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
 dihydroisocoumarinyl, isoindolinonyl,
 benzodioxanyl, benzoxazolinonyl, imidazopyrazolyl,
 quinazolinonyl, pyrazopyridyl, benzooxadiazolyl,
 15 dihydropyrimidinonyl, dihydrobenzofuranonyl,

where the R_C-heteroaryl group is bonded by any atom of the
 parent R_C-heteroaryl group substituted by hydrogen such that the new
 bond to the R_C-heteroaryl group replaces the hydrogen atom and its
 bond, where heteroaryl is optionally substituted 1, 2, 3, or 4
 20 groups that are independently:

(1) C₁-C₆ alkyl, optionally substituted with 1, 2, or
 3 groups independently selected from the group consisting of C₁-
 C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy,
 and -NR_{1-a}R_{1-b},

25 (2) -OH,
 (3) -NO₂,
 (4) -F, -Cl, -Br, -I,
 (5) -CO-OH,
 (6) -C≡N,

30 (V) C₂-C₁₀ alkenyl optionally substituted with one, two
 or three substituents independently selected from the group
 consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N,
 -CF₃, C₁-C₆ alkoxy, -O-phenyl, and -NR_{1-a}R_{1-b},

(VI) C₂-C₁₀ alkynyl optionally substituted with one, two or three substituents independently selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, and -NR_{1-a}R_{1-b},

5 (VII) -(C₁-C₆ alkyl)-O-(C₁-C₆ alkyl)-OH,

(VIII) -CH₂-NH-CH₂-CH(-O-CH₂-CH₃)₂,

(IX) -(CH₂)₀₋₆-C(=NR_{1-a})(NR_{1-a}R_{1-b});

where R_N is

(I) R_{N-1}-X_N- where X_N is -CO-, and where R_{N-1} is selected
10 from the group consisting of:

(A) phenyl, which is optionally substituted with one, two or three of the following substituents which can be the same or different and are:

(1) C₁-C₆ alkyl, optionally substituted with one,
15 two or three substituents selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

wherein R_{1-a} and R_{1-b} at each occurrence are independently H or C₁-C₆ alkyl,

20 (2) -OH,

(3) -NO₂,

(4) -F, -Cl, -Br, -I,

(5) -CO₂H,

(6) -C≡N,

25 (7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are the same or different and are selected from the group consisting of:

(a) -H,

(b) -C₁-C₈ alkyl optionally substituted with one substituent selected from the group consisting of:

30 (i) -OH,

(ii) -NH₂,

(iii) phenyl,

(c) -C₁-C₈ alkyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -Br, or -I,

- (d) -C₃-C₈ cycloalkyl,
(e) -(C₁-C₂ alkyl)-(C₃-C₈ cycloalkyl),
(f) -(C₁-C₆ alkyl)-O-(C₁-C₃ alkyl),
(g) -C₂-C₆ alkenyl,
5 (h) -C₂-C₆ alkynyl,
(i) -C₁-C₆ alkyl chain with one double bond
and one triple bond,
(j) -R_{1-aryl}, wherein R_{1-aryl} at each occurrence
is independently phenyl, naphthyl, indanyl, indenyl,
10 dihydronaphthyl, or tetralinyl each of which is optionally
substituted with 1, 2, 3, or 4 groups that are independently:
(i) C₁-C₆ alkyl optionally
substituted with one, two or three substituents independently
selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br,
15 -I, -OH, -SH, -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C₁-C₃ alkoxy,
(ii) C₂-C₆ alkenyl with one or two
double bonds, optionally substituted with one, two or three
substituents independently selected from the group consisting of
-F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},
20 (iii) C₂-C₆ alkynyl optionally
substituted with 1, 2, or 3 groups that are independently
selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -
CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},
(iv) -F, Cl, -Br and -I,
25 (v) -C₁-C₆ alkoxy optionally
substituted with 1, 2, or 3 -F,
(vi) -NR_{N-2}R_{N-3},
(vii) -OH,
(viii) -C≡N,
30 (ix) C₃-C₇ cycloalkyl, optionally
substituted with 1, 2, or 3 groups that are selected from the
group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy,
and -NR_{1-a}R_{1-b},
(x) -CO-(C₁-C₄ alkyl),

(xi) $-\text{SO}_2-\text{NR}_{1-a}\text{R}_{1-b}$,

(xii) $-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$, or

(xiii) $-\text{SO}_2-(\text{C}_1-\text{C}_4 \text{ alkyl})$,

(k) $-\text{R}_{1\text{-heteroaryl}}$, wherein $\text{R}_{1\text{-heteroaryl}}$ at each

5 occurrence is independently selected from the group consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, 10 indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, 15 isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl, 20 dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl, dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl, coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-oxide, tetrahydroquinolinyl, dihydroquinolinyl, dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl, 25 dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl, benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide, pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide, indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide, quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide, 30 thiazolyl N-oxide, indoliziny N-oxide, indazolyl N-oxide, benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and 35 benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(i) C_1-C_6 alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C_1-C_3 alkoxy,

(ii) C_2-C_6 alkenyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C_1-C_3 alkoxy, or -NR_{1-a}R_{1-b},

(iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C_1-C_3 alkoxy, or -NR_{1-a}R_{1-b},

(iv) -F, -Cl, -Br and -I,

(v) $-C_1-C_6$ alkoxy optionally substituted with one, two, or three -F,

(vi) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,

(vii) -OH,

(viii) -C≡N,

(ix) $(CH_2)_{0-4}-C_3-C_7$ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C_1-C_3 alkoxy, and -NR_{1-a}R_{1-b},

(x) $(CH_2)_{0-4}-CO-(C_1-C_6 \text{ alkyl})$,

(xi) $(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,

(xii) $(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,

(xiii) $(CH_2)_{0-4}-SO_2-(C_1-C_6 \text{ alkyl})$,

(xiv) $(CH_2)_{0-4}-N(R_{N-2})-SO_2-$, and

(xv) $(CH_2)_{0-4}-N(R_{N-2})-C(O)-$,

(1) $-R_1$ -heterocycle, wherein

R_1 -heterocycle at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl,

tetrahydrothienyl, homopiperidinyl, homomorpholinyl,
homothiomorpholinyl, homothiomorpholinyl S,S-dioxide,
oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl,
dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl,
5 dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide,
tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the $R_{1\text{-heterocycle}}$ group is bonded by
any atom of the parent $R_{1\text{-heterocycle}}$ group substituted by hydrogen
such that the new bond to the $R_{1\text{-heterocycle}}$ group replaces the
10 hydrogen atom and its bond, where heterocycle is optionally
substituted with 1, 2, 3, or 4 groups that are independently:

(a) $C_1\text{-}C_6$ alkyl optionally
substituted with one, two or three substituents independently
selected from the group consisting of $C_1\text{-}C_3$ alkyl, halogen, -OH,
15 -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and $C_1\text{-}C_3$ alkoxy,

(b) $C_2\text{-}C_6$ alkenyl with one or two
double bonds, optionally substituted with one, two or three
substituents independently selected from the group consisting of
-F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, $C_1\text{-}C_3$ alkoxy, and $-NR_{1-a}R_{1-b}$

20 (c) $C_2\text{-}C_6$ alkynyl with one or two
triple bonds, optionally substituted with one, two or three
substituents independently selected from the group consisting of
-F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, $C_1\text{-}C_3$ alkoxy, and $-NR_{1-a}R_{1-b}$

(d) halogen,

25 (e) $C_1\text{-}C_6$ alkoxy,
(f) $-C_1\text{-}C_6$ alkoxy optionally
substituted with one, two, or three -F,

(g) $-NR_{N-2}R_{N-3}$,

(h) -OH,

30 (i) $-C\equiv N$,
(j) $(CH_2)_{0-4}$ -($C_3\text{-}C_8$ cycloalkyl),
optionally substituted with 1, 2, or 3 groups independently
selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$,
 $-CF_3$, $C_1\text{-}C_3$ alkoxy, and $-NR_{1-a}R_{1-b}$,

- (k) - $(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_4 \text{ alkyl})$,
 (l) - $(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{1-a}\text{R}_{1-b}$,
 (m) - $(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$,
 (n) - $(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_6 \text{ alkyl})$, and
 (o) =O,
 (p) - $(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{N-2})-\text{SO}_2-$
 (q) - $(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{N-2})-\text{C}(\text{O})-$
- (8) - $(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
 (9) - $(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkenyl})$,
 (10) - $(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkynyl})$,
 (11) - $(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
 (12) - $(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-aryl}$,
 (13) - $(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl}$,
 (14) - $(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle}$,
 (15) - $(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{N-4}$ wherein R_{N-4} is selected from the group consisting of phenyl, morpholinyl, thiomorpholinyl, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl, thienyl, pyrazolyl, pyridyl N-oxide, oxazolyl, thiazolyl, imidazolyl, and pyrrolidinyl where each group is optionally substituted with one, two, three, or four groups that are independently C_1-C_6 alkyl,
- (16) - $(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{N-5}$ where R_{N-5} is selected from the group consisting of:
- (a) C_1-C_6 alkyl,
 (b) - $(\text{CH}_2)_{0-2}-(\text{R}_1\text{-aryl})$,
 (c) C_2-C_6 alkenyl,
 (d) C_2-C_6 alkynyl,
 (e) - $(\text{CH}_2)_{0-2}-\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,
 (f) - $(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heteroaryl})$, and
 (g) - $(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heterocycle})$,
- (17) - $(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{N-2}\text{R}_{N-3}$,
 (18) - $(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl})$,
 (19) - $(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
 (20) - $(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,

(21) - $(\text{CH}_2)_{0-4}\text{-N(H or R}_{N-5})\text{-CO-O-R}_{N-5}$,

(22) - $(\text{CH}_2)_{0-4}\text{-N(H or R}_{N-5})\text{-CO-N(R}_{N-5})_2$,

(23) - $(\text{CH}_2)_{0-4}\text{-N-CS-N(R}_{N-5})_2$,

(24) - $(\text{CH}_2)_{0-4}\text{-N(H or R}_{N-5})\text{-CO-R}_{N-2}$,

5 (25) - $(\text{CH}_2)_{0-4}\text{-NR}_{N-2}\text{R}_{N-3}$,

(26) - $(\text{CH}_2)_{0-4}\text{-R}_{N-4}$,

(27) - $(\text{CH}_2)_{0-4}\text{-O-CO-(C}_1\text{-C}_6\text{ alkyl)}$,

(28) - $(\text{CH}_2)_{0-4}\text{-O-P(O)-(OR}_{100})_2$ wherein

R_{100} at each occurrence is independently -H

10 or $\text{C}_1\text{-C}_4$ alkyl,

(29) - $(\text{CH}_2)_{0-4}\text{-O-CO-N(R}_{N-5})_2$,

(30) - $(\text{CH}_2)_{0-4}\text{-O-CS-N(R}_{N-5})_2$,

(31) - $(\text{CH}_2)_{0-4}\text{-O-(R}_{N-5})$,

(32) - $(\text{CH}_2)_{0-4}\text{-O-(R}_{N-5})\text{-COOH}$,

15 (33) - $(\text{CH}_2)_{0-4}\text{-S-(R}_{N-5})$,

(34) - $(\text{CH}_2)_{0-4}\text{-O-(C}_1\text{-C}_6\text{ alkyl optionally substituted with one, two, three, four, or five of -F)}$,

(35) $\text{C}_3\text{-C}_8$ cycloalkyl,

(36) $\text{C}_2\text{-C}_6$ alkenyl optionally substituted with $\text{C}_1\text{-}$

20 C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $\text{-C}\equiv\text{N}$, -CF_3 , $\text{C}_1\text{-C}_3$ alkoxy, or $\text{-NR}_{1-a}\text{R}_{1-b}$,

(37) $\text{C}_2\text{-C}_6$ alkynyl optionally substituted with $\text{C}_1\text{-}$
 C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $\text{-C}\equiv\text{N}$, -CF_3 , $\text{C}_1\text{-C}_3$ alkoxy, or $\text{-NR}_{1-a}\text{R}_{1-b}$,

25 (38) - $(\text{CH}_2)_{0-4}\text{-N(H or R}_{N-5})\text{-SO}_2\text{-R}_{N-2}$,

(39) - $(\text{CH}_2)_{1-4}\text{-(C}_3\text{-C}_8\text{ cycloalkyl)}$,

(B) $\text{-R}_{N\text{-heteroaryl}}$ where $\text{R}_{N\text{-heteroaryl}}$ is selected from the group consisting of pyridinyl, indolyl, indolinyl, isoindolyl, imidazolyl, isoxazolyl, oxazolyl, thiazolyl, indoliziny and
 30 isochromanyl,

where the $\text{R}_{N\text{-heteroaryl}}$ group is bonded by any atom of the parent $\text{R}_{N\text{-heteroaryl}}$ group substituted by hydrogen such that the new bond to the $\text{R}_{N\text{-heteroaryl}}$ group replaces the hydrogen atom and its bond, where heteroaryl is optionally substituted with one, two,
 35 three, or four of:

(1) C_1-C_6 alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

- 5 (2) -OH,
- (3) $-NO_2$,
- (4) -F, -Cl, -Br, -I,
- (5) $-CO_2H$,
- (6) $-C\equiv N$,
- 10 (7) $-(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,
- (8) $-(CH_2)_{0-4}-CO-(C_1-C_{12} \text{ alkyl})$,
- (9) $-(CH_2)_{0-4}-CO-(C_2-C_{12} \text{ alkenyl})$,
- (10) $-(CH_2)_{0-4}-CO-(C_2-C_{12} \text{ alkynyl})$,
- (11) $-(CH_2)_{0-4}-CO-(C_3-C_8 \text{ cycloalkyl})$,
- 15 (12) $-(CH_2)_{0-4}-CO-R_{1-aryl}$,
- (13) $-(CH_2)_{0-4}-CO-R_{1-heteroaryl}$,
- (14) $-(CH_2)_{0-4}-CO-R_{1-heterocycle}$,
- (15) $-(CH_2)_{0-4}-CO-R_{N-4}$,
- (16) $-(CH_2)_{0-4}-CO-O-R_{N-5}$,
- 20 (17) $-(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,
- (18) $-(CH_2)_{0-4}-SO-(C_1-C_8 \text{ alkyl})$,
- (19) $-(CH_2)_{0-4}-SO_2-(C_1-C_{12} \text{ alkyl})$,
- (20) $-(CH_2)_{0-4}-SO_2-(C_3-C_8 \text{ cycloalkyl})$,
- (21) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO-O-R_{N-5}$,
- 25 (22) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO-N(R_{N-5})_2$,
- (23) $-(CH_2)_{0-4}-N-CS-N(R_{N-5})_2$,
- (24) $-(CH_2)_{0-4}-N(-H \text{ or } R_{N-5})-CO-R_{N-2}$,
- (25) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,
- (26) $-(CH_2)_{0-4}-R_{N-4}$,
- 30 (27) $-(CH_2)_{0-4}-O-CO-(C_1-C_6 \text{ alkyl})$,
- (28) $-(CH_2)_{0-4}-O-P(O)-(OR_{100})_2$,
- (29) $-(CH_2)_{0-4}-O-CO-N(R_{N-5})_2$,
- (30) $-(CH_2)_{0-4}-O-CS-N(R_{N-5})_2$,
- (31) $-(CH_2)_{0-4}-O-(R_{N-5})$,
- 35 (32) $-(CH_2)_{0-4}-O-(R_{N-5})-COOH$,

(33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,

(34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl optionally substituted with one, two, three, four, or five of } -\text{F})$,

(35) $\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,

5 (36) $\text{C}_2-\text{C}_6 \text{ alkenyl optionally substituted with } \text{C}_1-\text{C}_3 \text{ alkyl, } -\text{F, } -\text{Cl, } -\text{Br, } -\text{I, } -\text{OH, } -\text{SH, } -\text{C}\equiv\text{N, } -\text{CF}_3, \text{ C}_1-\text{C}_3 \text{ alkoxy, or } -\text{NR}_{1-a}\text{R}_{1-b}$,

(37) $\text{C}_2-\text{C}_6 \text{ alkynyl optionally substituted with } \text{C}_1-\text{C}_3 \text{ alkyl, } -\text{F, } -\text{Cl, } -\text{Br, } -\text{I, } -\text{OH, } -\text{SH, } -\text{C}\equiv\text{N, } -\text{CF}_3, \text{ C}_1-\text{C}_3 \text{ alkoxy,}$
 10 or $-\text{NR}_{1-a}\text{R}_{1-b}$,

(38) $-(\text{CH}_2)_{0-4}-\text{N}(-\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$,

(39) $-(\text{CH}_2)_{1-4}-\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,

(C) $\text{R}_{\text{N-aryl}}-\text{W}-\text{R}_{\text{N-aryl}}$,

(D) $\text{R}_{\text{N-aryl}}-\text{W}-\text{R}_{\text{N-heteroaryl}}$,

15 (E) $\text{R}_{\text{N-aryl}}-\text{W}-\text{R}_{1-\text{heterocycle}}$,

(F) $\text{R}_{\text{N-heteroaryl}}-\text{W}-\text{R}_{\text{N-aryl}}$,

(G) $\text{R}_{\text{N-heteroaryl}}-\text{W}-\text{R}_{\text{N-heteroaryl}}$,

(H) $\text{R}_{\text{N-heteroaryl}}-\text{W}-\text{R}_{\text{N-1-heterocycle}}$,

(I) $\text{R}_{\text{N-heterocycle}}-\text{W}-\text{R}_{\text{N-aryl}}$,

20 (J) $\text{R}_{\text{N-heterocycle}}-\text{W}-\text{R}_{\text{N-heteroaryl}}$,

(K) $\text{R}_{\text{N-heterocycle}}-\text{W}-\text{R}_{\text{N-1-heterocycle}}$,

where W is

(19) $-(\text{CH}_2)_{1-4}-$,

(20) $-\text{O}-$,

25 (21) $-\text{S}(\text{O})_{0-2}-$,

(22) $-\text{N}(\text{R}_{\text{N}-5})-$,

(23) $-\text{CO}-$; or

(24) a bond;

(II) $-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})-\text{M}-(\text{C}_1-\text{C}_6 \text{ alkyl})$, where M is S, SO or
 30 SO_2 , and wherein each alkyl is unsubstituted or substituted with one, two, or three of substituents independently selected from the group consisting of:

(A) $-\text{NH}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,

(B) $-\text{NH}-\text{CO}-\text{O}-\text{R}_{\text{N-8}}$,

35 (C) $-\text{NR}_{\text{N-2}}\text{R}_{\text{N-3}}$;

where R_1 is

$-(CH_2)_{n_1}$ -phenyl, where n_1 is zero or one, and which is optionally substituted with one, two, three or four of the following substituents on the phenyl ring:

5 (A) C_1 - C_6 alkyl optionally substituted with one, two or three substituents selected from the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF $_3$, C_1 - C_3 alkoxy, and -NR $_{1-a}$ R $_{1-b}$,

10 (B) C_2 - C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF $_3$, C_1 - C_3 alkoxy, and -NR $_{1-a}$ R $_{1-b}$,

15 (C) C_2 - C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF $_3$, C_1 - C_3 alkoxy, and -NR $_{1-a}$ R $_{1-b}$,

(D) -F, Cl, -Br or -I,

(F) $-C_1$ - C_6 alkoxy optionally substituted with one, two or three of -F,

20 (G) -NR $_{N-2}$ R $_{N-3}$ where R $_{N-2}$ and R $_{N-3}$ are as defined below,

(H) -OH,

(I) -C \equiv N,

25 (J) C_3 - C_7 cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF $_3$, C_1 - C_3 alkoxy, and -NR $_{1-a}$ R $_{1-b}$,

(K) -CO-(C_1 - C_4 alkyl),

(L) -SO $_2$ -NR $_{1-a}$ R $_{1-b}$,

30 (M) -CO-NR $_{1-a}$ R $_{1-b}$,

(N) -SO $_2$ -(C_1 - C_4 alkyl); and

where R_2 is

(I) -(Z)- C_1 - C_6 alkyl, where Z is a bond, -C(O)-, -CO $_2$ - or -SO $_2$ -, wherein the alkyl group is optionally substituted with

one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, C₁-C₇ alkyl (optionally substituted with C₁-C₃ alkyl and C₁-C₃ alkoxy), -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, -NR_{1-a}R_{1-b} where R_{1-a} and R_{1-b} are
5 independently -H or C₁-C₆ alkyl, and -OC=O NR_{1-a}R_{1-b},

(II) -(Z)-CH₂-S(O)₀₋₂-(C₁-C₆ alkyl),

(III) -(Z)-CH₂-CH₂-S(O)₀₋₂-(C₁-C₆ alkyl),

(IV) -(Z)-C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents
10 selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(V) -(Z)-C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -
15 CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(VI) -(Z)-(CH₂)_{n1}-(R_{1-aryl}), where Z is a bond, CO, CO₂ or SO₂, where n₁ is zero or one and where R_{1-aryl} is phenyl, 1-naphthyl, 2-naphthyl and indanyl, indenyl, dihydronaphthalyl, or tetralinyl optionally substituted with one, two, three or four
20 of the following substituents on the aryl ring:

(A) C₁-C₆ alkyl optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

25 (B) C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(C) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents
30 selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(D) -F, Cl, -Br or -I,

(F) $-C_1-C_6$ alkoxy optionally substituted with one, two or three of - F,

(G) $-NR_{N-2}R_{N-3}$ where R_{N-2} and R_{N-3} are as defined below,

5 (H) $-OH$,

(I) $-C\equiv N$,

(J) C_3-C_7 cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and -

10 $NR_{1-a}R_{1-b}$,

(K) $-CO-(C_1-C_4 \text{ alkyl})$,

(L) $-SO_2-NR_{1-a}R_{1-b}$,

(M) $-CO-NR_{1-a}R_{1-b}$,

(N) $-SO_2-(C_1-C_4 \text{ alkyl})$,

15 (VII) $-(Z)-(CH_2)_{n_1}-(R_{1-\text{heteroaryl}})$ where n_1 is as defined above and where $R_{1-\text{heteroaryl}}$ is selected from the group consisting of:

pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, 20 quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, 25 isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanly, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranly, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranly, benzotetrahydrothienyl, 30 purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl, dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl, dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl, coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N- 35 oxide, tetrahydroquinolinyl, dihydroquinolinyl,

dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl,
benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,
5 indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide,
quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-
oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
thiazolyl N-oxide, indolizinyll N-oxide, indazolyl N-oxide,
benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
10 oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide,
benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is bonded to $-(CH_2)_{n1}-$ by any ring atom
of the parent R_N -heteroaryl group substituted by hydrogen such that
15 the new bond to the R_1 -heteroaryl group replaces the hydrogen atom
and its bond, where heteroaryl is optionally substituted with
one, two, three or four of:

(1) C_1-C_6 alkyl optionally substituted with one,
two or three substituents selected from the group consisting of
20 C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$,
 $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(2) C_2-C_6 alkenyl with one or two double bonds,
optionally substituted with one, two or three substituents
selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-$
25 CF_3 , C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(3) C_2-C_6 alkynyl with one or two triple bonds,
optionally substituted with one, two or three substituents
selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-$
30 CF_3 , C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(4) $-F$, Cl , $-Br$ or $-I$,

(6) $-C_1-C_6$ alkoxy optionally substituted with
one, two, or three of $-F$,

(7) $-NR_{N-2}R_{N-3}$ where R_{N-2} and R_{N-3} are as defined
below,

(8) -OH,

(9) -C≡N,

(10) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(11) -CO-(C₁-C₄ alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

(13) -CO-NR_{1-a}R_{1-b}, or

10 (14) -SO₂-(C₁-C₄ alkyl), with the proviso that when n₁ is zero R_{1-heteroaryl} is not bonded to the carbon chain by nitrogen, or

(VIII) -(Z)-(CH₂)_{n1}-(R_{1-heterocycle}) where n₁ is as defined above and R_{1-heterocycle} is selected from the group consisting of:
 15 morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl dihydropyrazinyl dihydropyridinyl dihydropyrimidinyl, dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, homothiomorpholinyl S-oxide,

25 where the R_{1-heterocycle} group is bonded by any atom of the parent R_{1-heterocycle} group substituted by hydrogen such that the new bond to the R_{1-heterocycle} group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with one, two, 30 three or four:

(1) C₁-C₆ alkyl optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(2) C₂-C₆ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

5 (3) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(4) -F, Cl, -Br, or -I,

10 (5) C₁-C₆ alkoxy,

(6) -C₁-C₆ alkoxy optionally substituted with one, two, or three -F,

(7) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

15 (8) -OH,

(9) -C≡N,

(10) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(11) -CO-(C₁-C₄ alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

(13) -CO-NR_{1-a}R_{1-b},

(14) -SO₂-(C₁-C₄ alkyl),

25 (15) =O, with the proviso that when n₁ is zero R_{1-heterocycle} is not bonded to the carbon chain by nitrogen; and

where R₂₀ is H or C₁₋₆ alkyl or alkenyl.

30 2. A compound according to claim 1, wherein R_c is -(CR_{c-x}R_{c-y})₀₋₄-R_{c-aryl} where R_{c-x} and R_{c-y} are independently selected from the group consisting of

-H,

C₁-C₄ alkyl optionally substituted with 1 or 2 -OH,

C₁-C₄ alkoxy optionally substituted with 1, 2, or 3
halogen,

-(CH₂)₀₋₄-C₃-C₈ cycloalkyl,

C₂-C₆ alkenyl containing one or two double bonds,

5 C₂-C₆ alkynyl containing one or two triple bonds, and
phenyl,

or

R_{C-x} and R_{C-y} are taken together with the carbon to
which they are attached to form a carbocycle of three, four,
10 five, six or seven carbon atoms, where one carbon atom is
optionally replaced by a group selected from -O-, -S-, -SO₂-,
-NR_{N-2}- and R_{C-aryl}, wherein

R_{C-aryl} is phenyl, which is optionally substituted
with 1, 2, or 3 groups that are independently:

15 (1) C₁-C₆ alkyl, optionally substituted with
one, two or three substituents selected from the group
consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃
alkoxy, and -NR_{1-a}R_{1-b},

(2) -OH,

20 (3) -NO₂,

(4) halogen,

(5) -CO₂H,

(6) -C≡N.

25 3. A compound according to claim 1, wherein R_C is -(CR_{C-x}
R_{C-y})₀₋₄-R_{C-aryl} where R_{C-x} and R_{C-y} are independently selected from
the group consisting of

-H,

C₁-C₄ alkyl optionally substituted with 1 or 2 -OH,

30 C₁-C₄ alkoxy optionally substituted with 1, 2, or 3
halogen,

-(CH₂)₀₋₄-C₃-C₈ cycloalkyl,

C₂-C₆ alkenyl containing one or two double bonds,

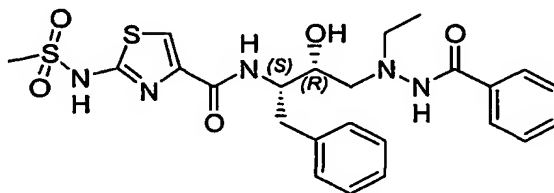
35 C₂-C₆ alkynyl containing one or two triple bonds, and
phenyl,

or

R_{C-x} and R_{C-y} are taken together with the carbon to which they are attached to form a carbocycle of three, four, five, six or seven carbon atoms, where one carbon atom is optionally replaced by a group selected from -O-, -S-, -SO₂-, -NR_{N-2}- and R_{C-aryl} , wherein

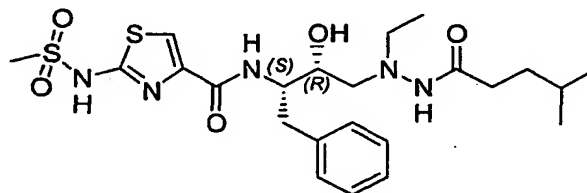
- (CR_{C-x}R_{C-y})₀₋₄-R_{C-heteroaryl} is selected from the group consisting of pyridinyl, indolyl, indolinyl, isoindolyl, imidazolyl, isoxazolyl, oxazolyl, thiazolyl, indolizinyll and isochromanyll.

4. A compound according to claim 1, of the formula:



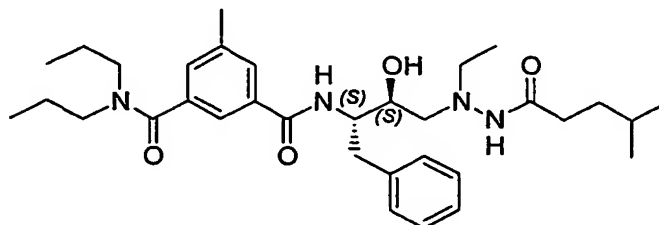
15

5. A compound according to claim 1, of the formula:

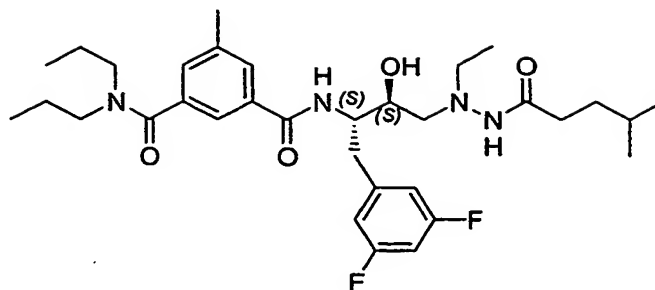


20

6. A compound according to claim 1, of the formula:

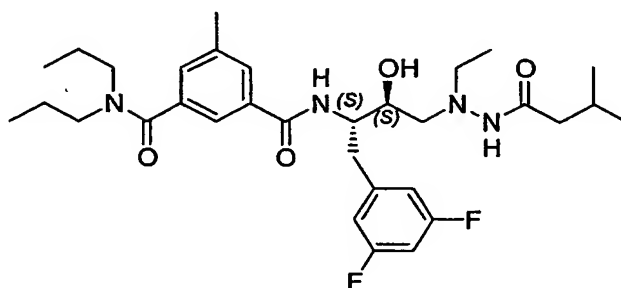


7. A compound according to claim 1, of the formula:



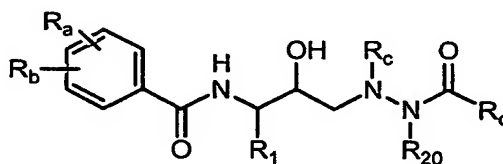
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8. A compound according to claim 1, of the formula:



10

9. A compound of the formula III:



(III)

or a pharmaceutically acceptable salt thereof wherein

15 R_1 represents phenyl (C_1 - C_6)alkyl where the phenyl is optionally substituted with up to three groups independently selected from halogen, hydroxy, C_1 - C_2 alkyl, C_1 - C_2 alkoxy, amino, nitro, trifluoromethyl, cyano, mono(C_1 - C_2)alkylamino and di(C_1 - C_2)alkylamino;

R_a and R_b independently represent hydrogen, hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, C_3 - C_7 cycloalkyl, C_3 - C_7 cycloalkyl(C_1 - C_6)alkyl, C_3 - C_7 cycloalkyl(C_1 - C_6)alkoxy, halogen, cyano, amino, mono(C_1 - C_6)alkylamino, di(C_1 - C_6)alkylamino, mono- or di(C_1 - C_6)alkylaminosulfonyl, C_1 - C_6 alkyl sulfonylamino, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, trifluoromethyl, mono(C_1 - C_6)alkylaminocarbonyl, or di(C_1 - C_6)alkylaminocarbonyl and provided that not both R_a and R_b are hydrogen simultaneously;

10 R_c represents hydrogen, or C_1 - C_6 alkyl, C_2 - C_6 alkenyl, or C_2 - C_6 alkynyl each of which is optionally substituted with halogen, hydroxy, amino, cyano, or trifluoromethyl;

R_d represents

phenyl optionally substituted with hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, C_3 - C_7 cycloalkyl, C_3 - C_7 cycloalkyl(C_1 - C_6)alkyl, C_3 - C_7 cycloalkyl(C_1 - C_6)alkoxy, halogen, cyano, amino, mono(C_1 - C_6)alkylamino, di(C_1 - C_6)alkylamino, mono- or di(C_1 - C_6)alkylaminosulfonyl, C_1 - C_6 alkyl sulfonylamino, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl; or

C_1 - C_6 alkyl optionally substituted with hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halogen, cyano, amino, mono(C_1 - C_6)alkylamino, di(C_1 - C_6)alkylamino, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, or trifluoromethyl; and

25 R_{20} represents hydrogen, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, or trifluoromethyl.

10. A compound according to claim 9, wherein R_1 is benzyl where the phenyl is optionally substituted.

5 11. A compound according to claim 10, wherein the phenyl is substituted with one or two groups independently selected from halogen, hydroxy, C_1 - C_3 alkyl, amino, and trifluoromethyl.

12. A compound according to claim 11, wherein phenyl is
10 substituted with two groups independently selected from halogen, hydroxy, and trifluoromethyl.

13. A compound according to claim 12, wherein phenyl is disubstituted with halogen.

15

14. A compound according to claim 13, wherein R_1 is 3,5-difluorobenzyl.

15. A compound according to claim 12, wherein R_a and R_b
20 are different and R_b represents mono- or di(C_1 - C_6)alkylaminocarbonyl.

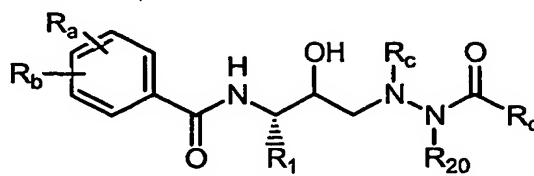
16. A compound according to claim 12, wherein R_d is phenyl optionally substituted with C_1 - C_3 alkyl, C_1 - C_3 alkoxy, amino,
25 hydroxy, or halogen.

17. A compound according to claim 12, wherein R_c is hydrogen or C_1 - C_4 alkyl.

18. A compound according to claim 17, wherein R_c is C_1 - C_3 alkyl.

19. A compound according to claim 12, wherein R_d is C_1 - C_6 lower alkyl and R_{20} is hydrogen.

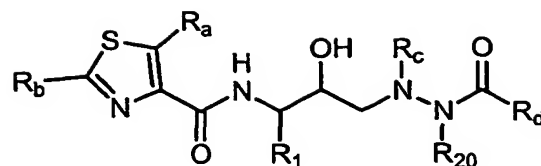
20. A compound according to claim 12, which has the formula IV:



(IV)

21. A compound according to claim 20, wherein R_1 is benzyl where the phenyl is disubstituted with chloro or fluoro;
 R_c is C_1 - C_3 alkyl;
 R_d is C_1 - C_6 lower alkyl;
 R_{20} is hydrogen or C_1 - C_6 alkyl; and
 R_b is di(C_1 - C_6)alkylaminocarbonyl attached to the 3-position of the phenyl group.

22. A compound of the formula V:



(V)

or a pharmaceutically acceptable salt thereof wherein

R₁ represents phenyl (C₁-C₆)alkyl where the phenyl is optionally
 5 substituted with up to three groups independently selected
 from halogen, hydroxy, C₁-C₂ alkyl, C₁-C₂ alkoxy, amino,
 nitro, trifluoromethyl, cyano, mono(C₁-C₂)alkylamino and
 di(C₁-C₂)alkylamino;

R_a and R_b independently represent hydrogen, hydroxy, C₁-C₆ alkyl,
 10 C₁-C₆ alkoxy, C₃-C₇ cycloalkyl, C₃-C₇ cycloalkyl(C₁-C₆)alkyl,
 C₃-C₇ cycloalkyl(C₁-C₆)alkoxy, halogen, cyano, amino,
 mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, mono- or di(C₁-
 C₆)alkylaminosulfonyl, C₁-C₆ alkyl sulfonylamino, C₂-C₆
 alkenyl, C₂-C₆ alkynyl, trifluoromethyl, mono(C₁-
 15 C₆)alkylaminocarbonyl, or di(C₁-C₆)alkylaminocarbonyl and
 provided that not both R_a and R_b are hydrogen
 simultaneously;

R_c represents hydrogen, or C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆
 alkynyl each of which is optionally substituted with
 20 halogen, hydroxy, amino, cyano, or trifluoromethyl;

R_d represents

phenyl optionally substituted with hydroxy, C₁-C₆ alkyl, C₁-
 C₆ alkoxy, C₃-C₇ cycloalkyl, C₃-C₇ cycloalkyl(C₁-
 C₆)alkyl, C₃-C₇ cycloalkyl(C₁-C₆)alkoxy, halogen,

cyano, amino, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, mono- or di(C₁-C₆)alkylaminosulfonyl, C₁-C₆ alkyl sulfonylamino, C₂-C₆ alkenyl, C₂-C₆ alkynyl; or

5 C₁-C₆ alkyl optionally substituted with hydroxy, C₁-C₆ alkyl, C₁-C₆ alkoxy, halogen, cyano, amino, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, C₂-C₆ alkenyl, C₂-C₆ alkynyl, or trifluoromethyl; and

R₂₀ represents hydrogen, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, or trifluoromethyl.

23. A compound according to claim 22, wherein R₁ is benzyl where the phenyl is optionally substituted.

15 24. A compound according to claim 23, wherein the phenyl is substituted with one or two groups independently selected from halogen, hydroxy, C₁-C₃ alkyl, amino, and trifluoromethyl.

25. A compound according to claim 24, wherein phenyl is substituted with two groups independently selected from halogen, hydroxy, and trifluoromethyl.

26. A compound according to claim 25, wherein phenyl is disubstituted with halogen.

27. A compound according to claim 26, wherein R₁ is 3,5-difluorobenzyl.

28. A compound according to claim 25, wherein R_a and R_b are different and R_b represents C₁-C₆)alkylsulfonylamino.

29. A compound according to claim 25, wherein R_d is phenyl optionally substituted with C₁-C₃ alkyl, C₁-C₃ alkoxy, amino, hydroxy, or halogen.

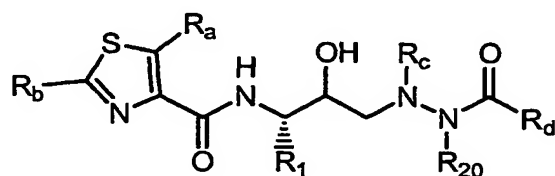
10

30. A compound according to claim 25, wherein R_c is hydrogen or C₁-C₄ alkyl.

31. A compound according to claim 30, wherein R_c is C₁-C₃ alkyl.

32. A compound according to claim 25, wherein R_d is C₁-C₆ lower alkyl and R₂₀ is hydrogen.

33. A compound according to claim 25, which has the formula VI:



(VI)

34. A compound according to claim 33, wherein

R_1 is benzyl where the phenyl is disubstituted with chloro or fluoro;

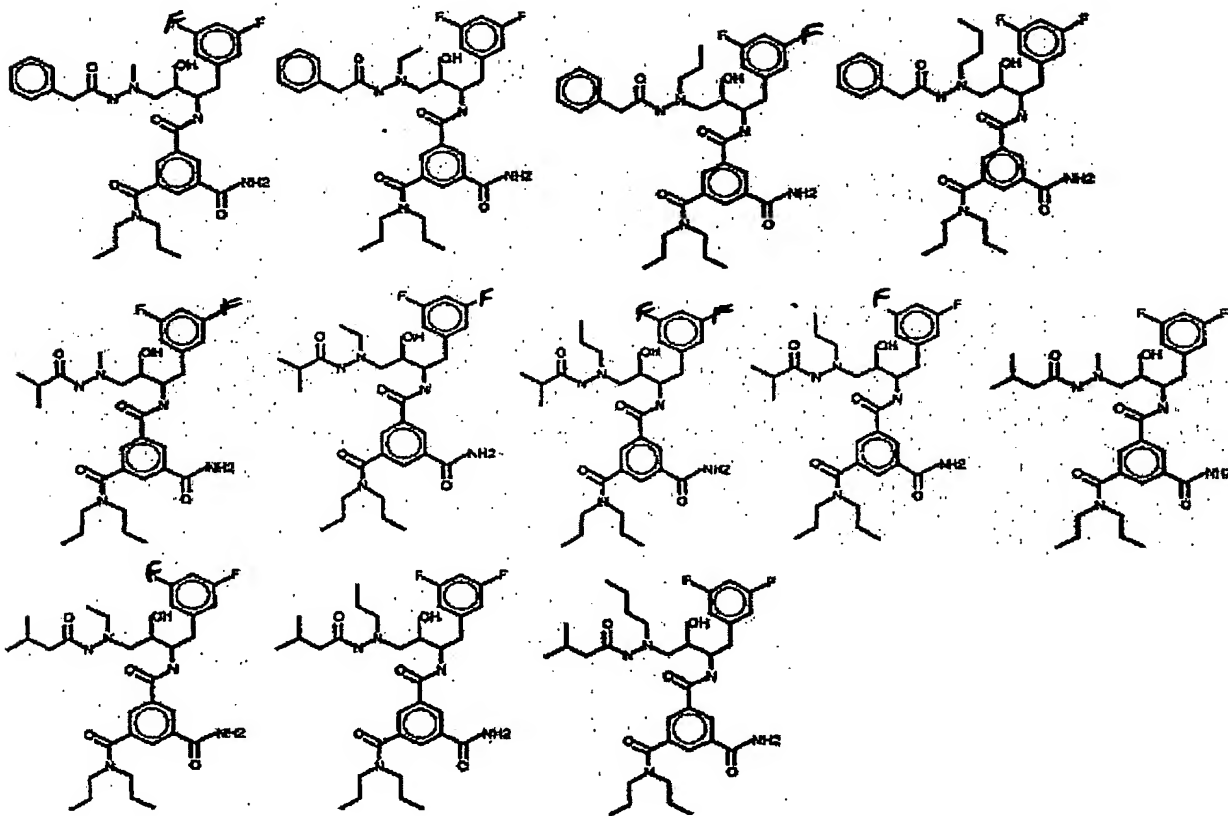
R_c is C_1 - C_3 alkyl;

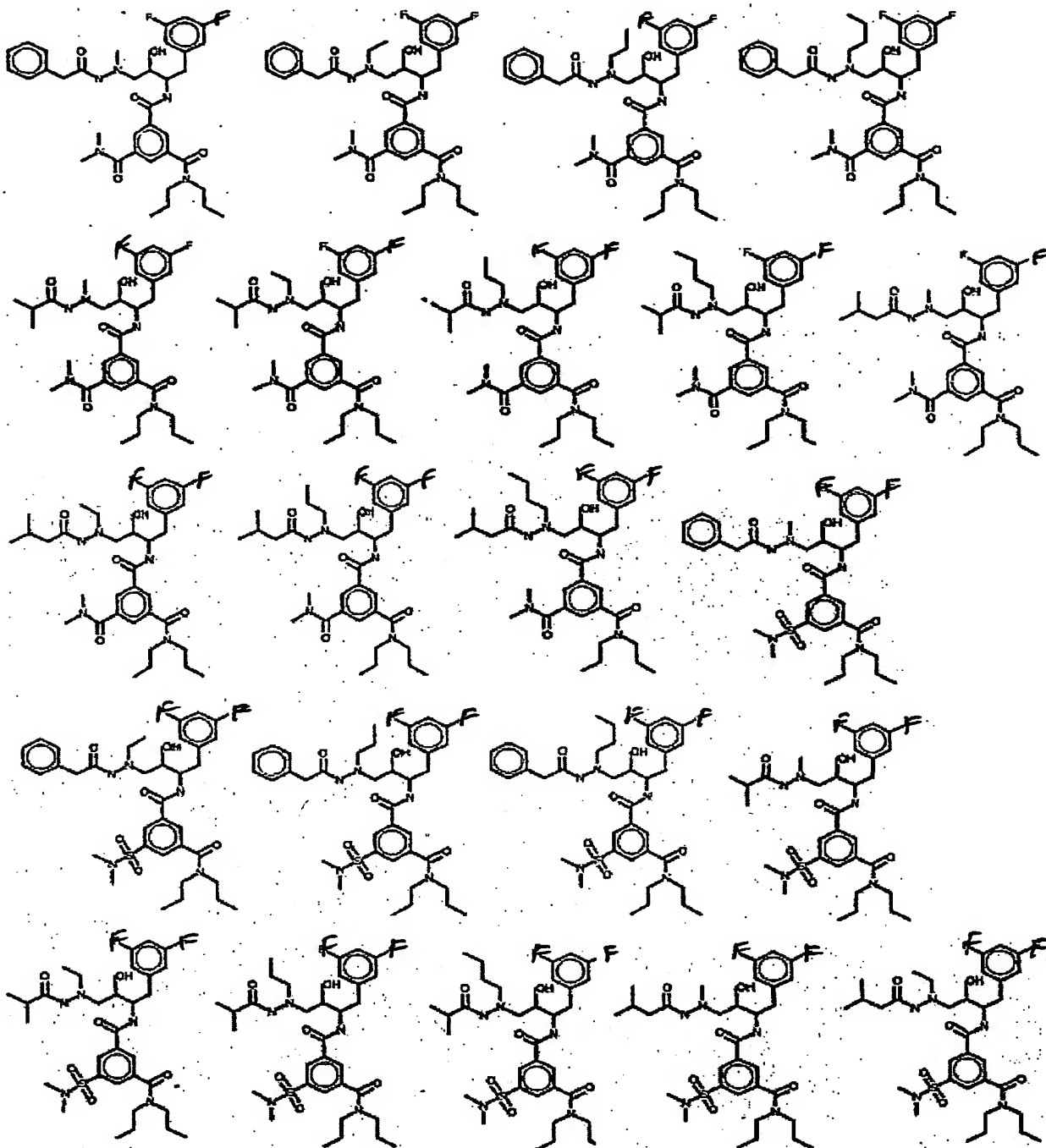
5 R_d is C_1 - C_6 lower alkyl;

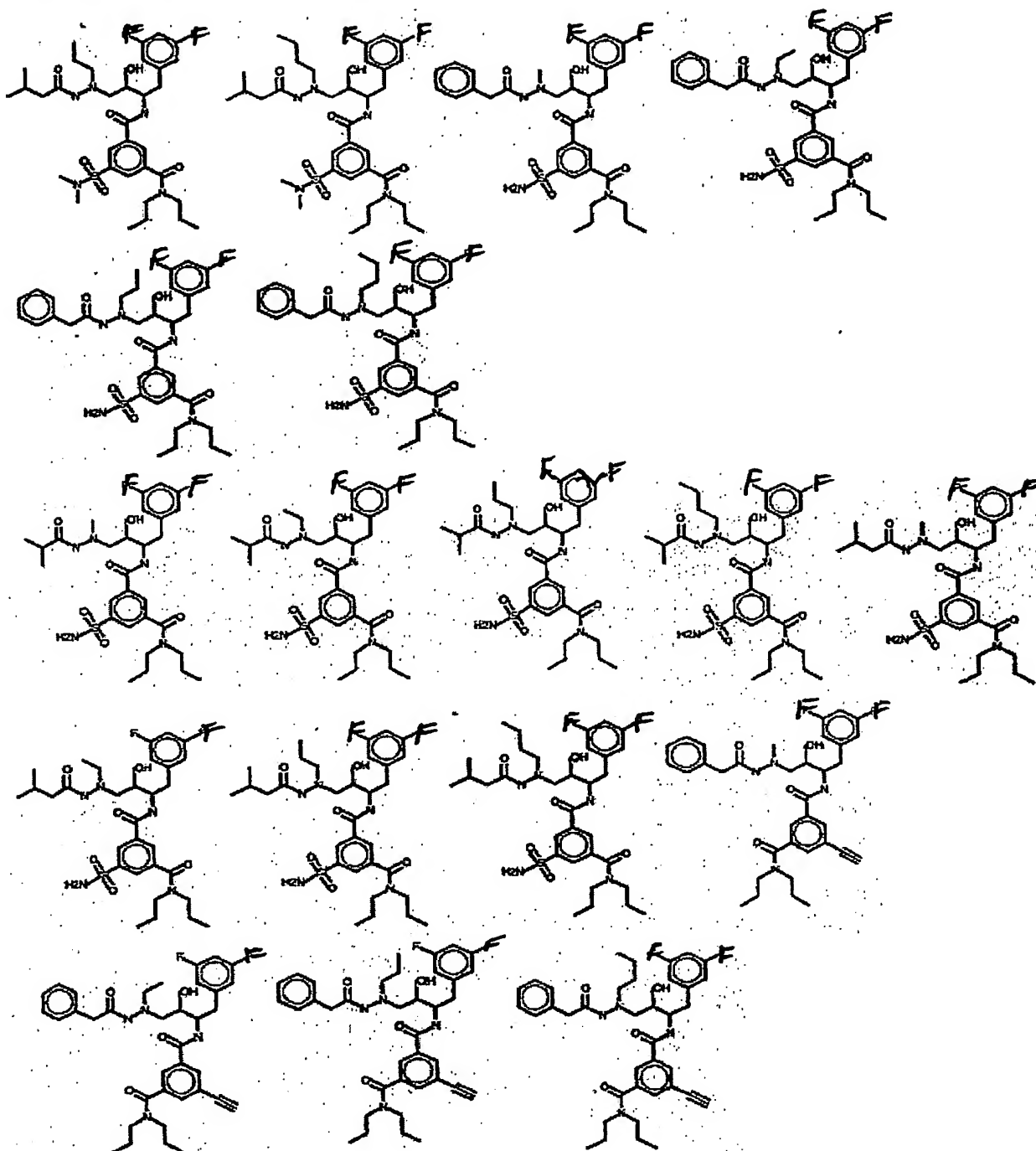
R_{20} is hydrogen or C_1 - C_6 alkyl; and

R_b is alkylsulfonylamino attached to the 2-position of the thiazolyl group.

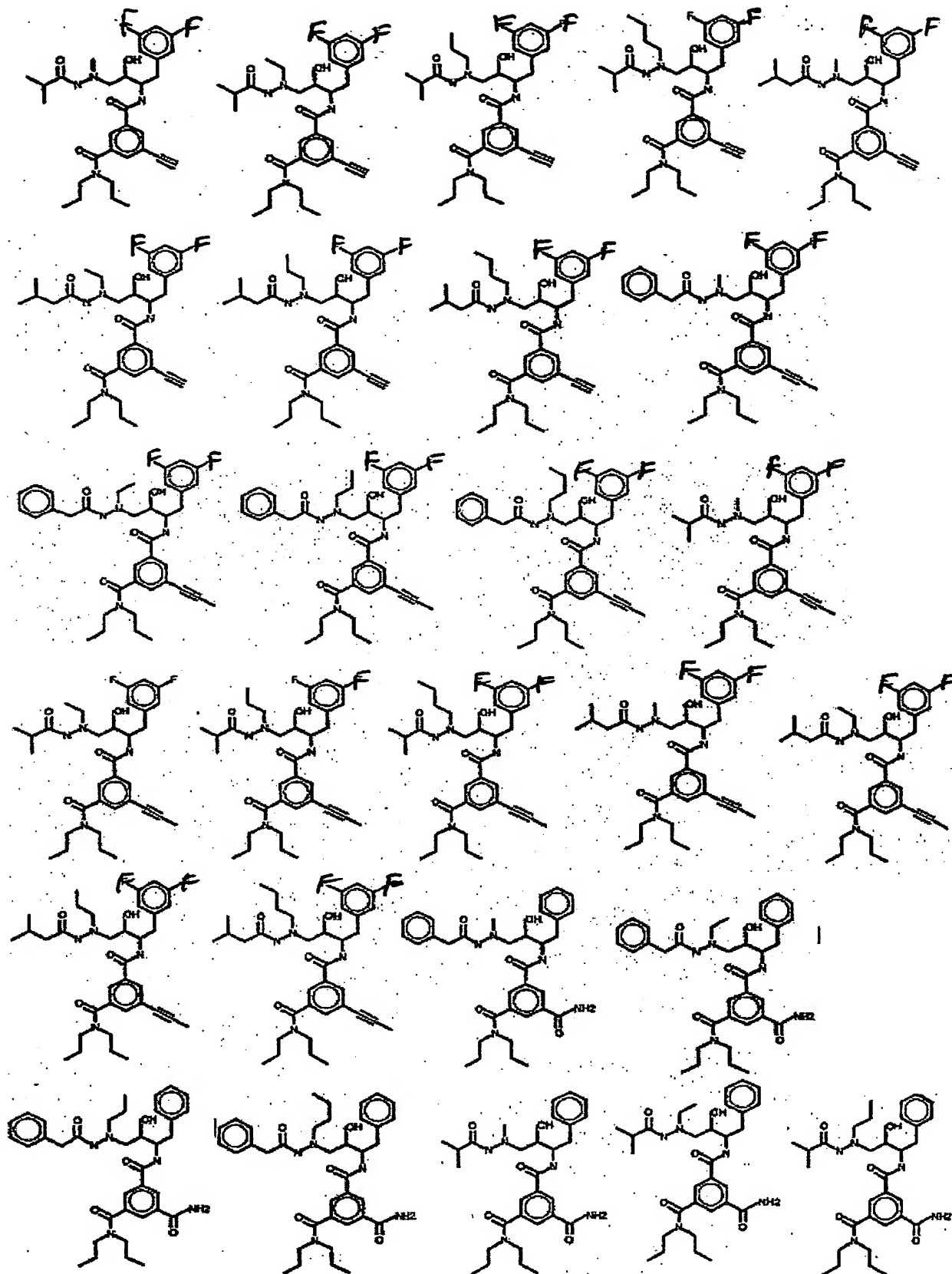
10 35. A compound according to claim 1, selected from the group consisting of:

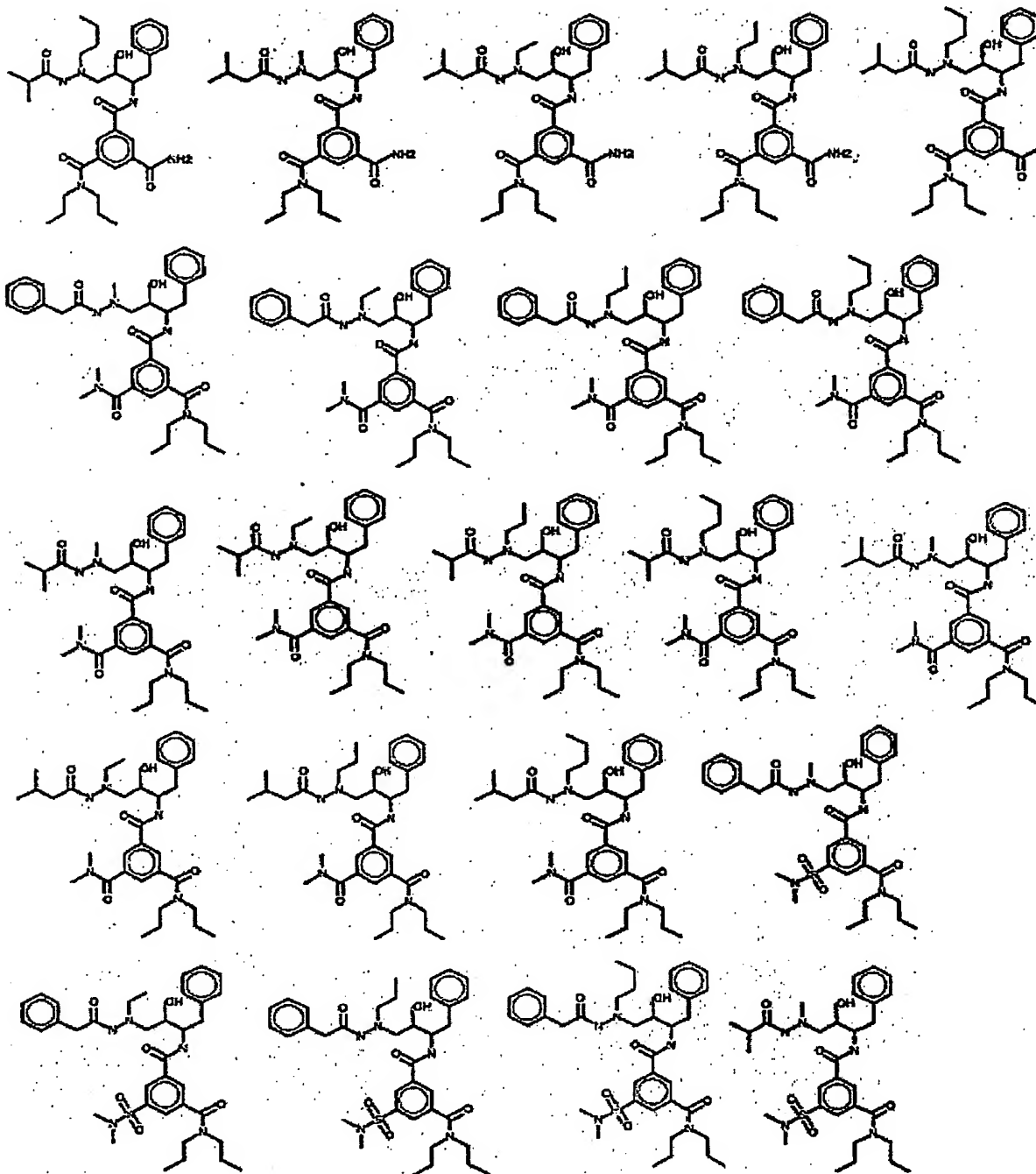


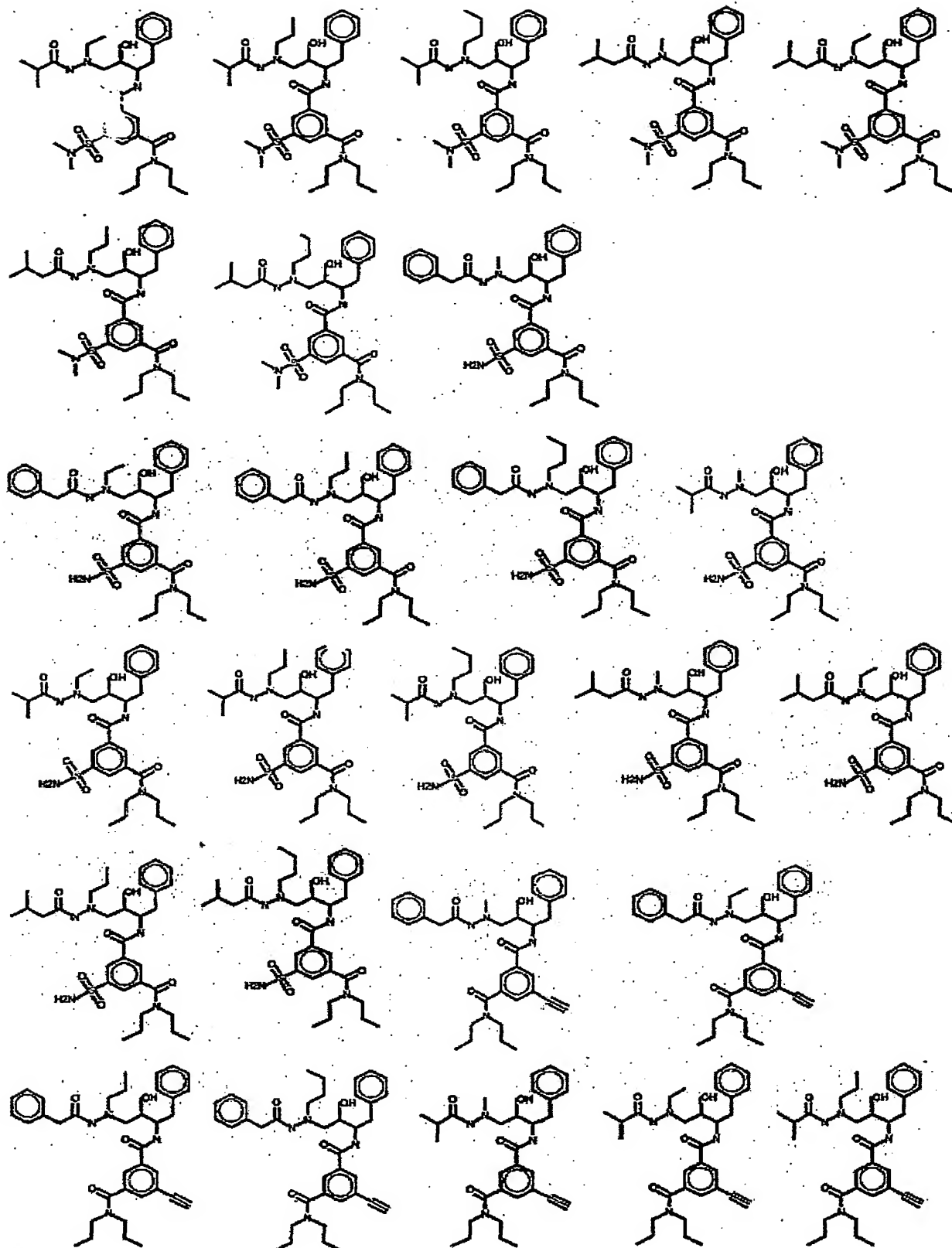


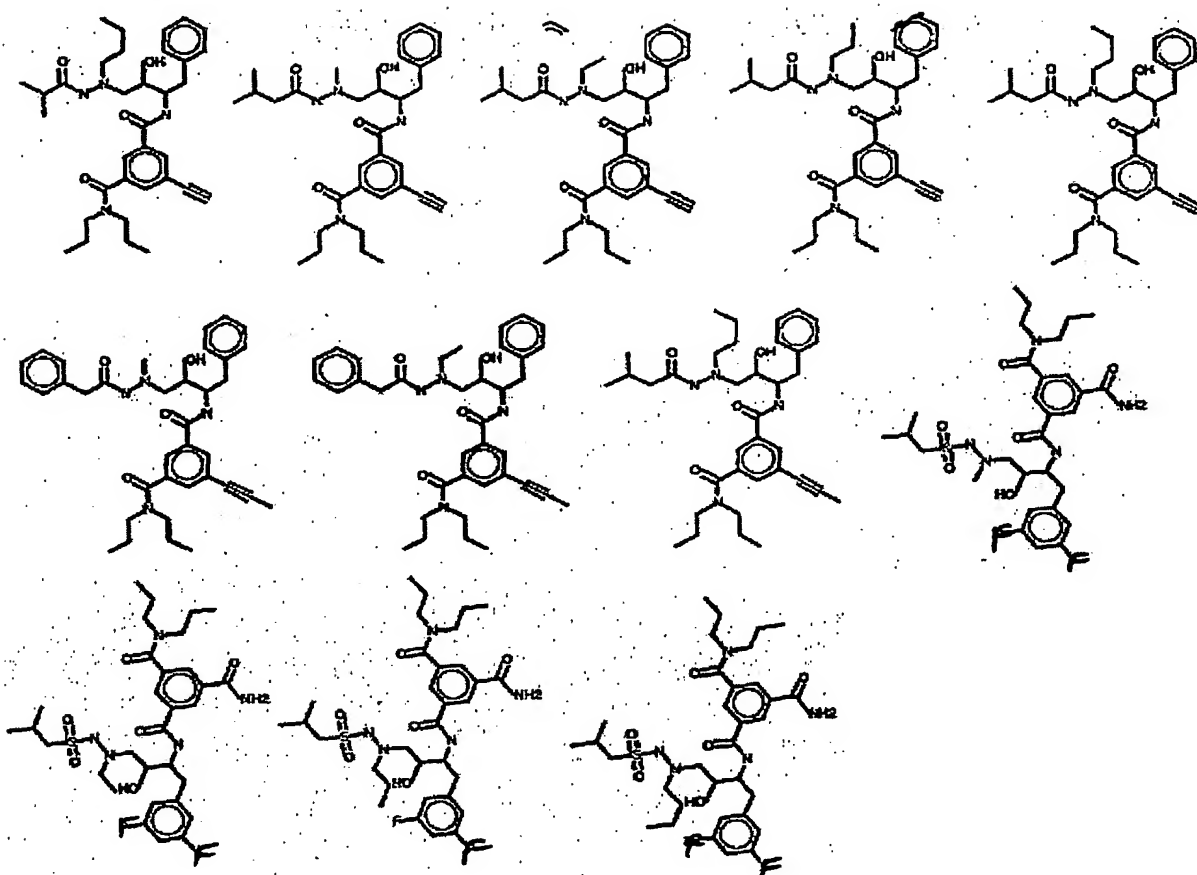


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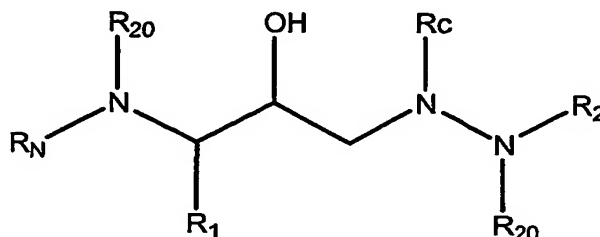


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36. A method of treating a patient who has, or in preventing a patient from getting, a disease or condition selected from the group consisting of Alzheimer's disease, for helping prevent or delay the onset of Alzheimer's disease, for treating patients with mild cognitive impairment (MCI) and preventing or delaying the onset of Alzheimer's disease in those who would progress from MCI to AD, for treating Down's syndrome, for treating humans who have Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, for treating cerebral amyloid angiopathy and preventing its potential consequences, i.e. single and recurrent lobar hemorrhages, for treating other degenerative dementias, including dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's disease, dementia associated with progressive supranuclear palsy, dementia associated with cortical basal degeneration, diffuse Lewy body type of Alzheimer's disease and who is in need

of such treatment which comprises administration of a therapeutically effective amount of a compound selected from the group consisting of an aza hydroxylated ethyl amine of the formula II:

5



or a pharmaceutically acceptable salt thereof,

10 where R_c is

(I) -C₁-C₁₀ alkyl optionally substituted with one, two or three groups independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, -NR_{1-a}R_{1-b}, -OC(=O)NR_{1-a}R_{1-b}, -S(=O)₀₋₂R_{1-a}, -NR_{1-a}C(=O)NR_{1-a}R_{1-b},
15 -C(=O)NR_{1-a}R_{1-b}, and -S(=O)₂NR_{1-a}R_{1-b} wherein

R_{1-a} and R_{1-b} at each occurrence are independently H or C₁-C₆ alkyl,

(II) -(CH₂)₀₋₃-(C₃-C₈) cycloalkyl where cycloalkyl can be optionally substituted with one, two or three substituents
20 independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, -O-phenyl, -CO₂H, -CO₂-(C₁-C₄ alkyl), and -NR_{1-a}R_{1-b}

(III) -(CR_{C-x}R_{C-y})₀₋₄-R_{C-aryl} where R_{C-x} and R_{C-y} are independently selected from the group consisting of

25 -H,
C₁-C₄ alkyl optionally substituted with 1 or 2 -OH,
C₁-C₄ alkoxy optionally substituted with 1, 2, or 3
halogen,
-(CH₂)₀₋₄-C₃-C₈ cycloalkyl,
30 C₂-C₆ alkenyl containing one or two double bonds,
C₂-C₆ alkynyl containing one or two triple bonds, and

phenyl,

or

R_{C-x} and R_{C-y} are taken together with the carbon to which they are attached to form a carbocycle of three, four, five, six or seven carbon atoms, where one carbon atom is optionally replaced by a group selected from -O-, -S-, -SO₂-, -NR_{N-2}- and R_{C-aryl} , wherein

R_{C-aryl} is phenyl, which is optionally substituted with 1, 2, or 3 groups that are independently:

(1) C₁-C₆ alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(2) -OH,

(3) -NO₂,

(4) halogen,

(5) -CO₂H,

(6) -C≡N,

(7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3} where

R_{N-2} and R_{N-3} are independently selected from the group consisting of:

(a) -H,

(b) -C₁-C₆ alkyl optionally substituted with one substituent selected from the group consisting of:

(i) -OH, and

(ii) -NH₂,

(c) -C₁-C₆ alkyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -Br, -I, or OH,

(d) -C₃-C₇ cycloalkyl,

(e) -(C₁-C₂ alkyl)-(C₃-C₇ cycloalkyl),

(f) -(C₁-C₆ alkyl)-O-(C₁-C₃ alkyl),

(g) -C₂-C₆ alkenyl

(h) -C₂-C₆ alkynyl

(i) $-C_1-C_6$ alkyl chain with one double bond and one triple bond,

(j) $-R_{1-aryl}$ wherein R_{1-aryl} at each occurrence is independently phenyl, naphthyl, indanyl, indenyl, dihydronaphthyl, or tetralinyl each of which is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(i) C_1-C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

(ii) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(iv) $-F$, Cl , $-Br$ and $-I$,

(v) $-C_1-C_6$ alkoxy optionally substituted with 1, 2, or 3 $-F$,

(vi) $-NR_{N-2}R_{N-3}$,

(vii) $-OH$,

(viii) $-C\equiv N$,

(ix) C_3-C_7 cycloalkyl, optionally substituted with 1, 2, or 3 groups that are selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(x) $-CO-(C_1-C_4 \text{ alkyl})$,

(xi) $-SO_2-NR_{1-a}R_{1-b}$,

(xii) $-CO-NR_{1-a}R_{1-b}$, or

(xiii) $-SO_2-(C_1-C_4 \text{ alkyl})$,

(k) $-R_{1-heteroaryl}$ wherein $R_{1-heteroaryl}$ at each occurrence is independently selected from the group

consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indolizinyl, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl, dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl, dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl, coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-oxide, tetrahydroquinolinyl, dihydroquinolinyl, dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl, dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl, benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide, pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide, indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide, quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide, thiazolyl N-oxide, indolizinyl N-oxide, indazolyl N-oxide, benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(i) C_1-C_6 alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -NR_{1-a}R_{1-b}, -C≡N, -CF₃, and C_1-C_3 alkoxy,

5 (ii) C_2-C_6 alkenyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, -C≡N, -CF₃, C_1-C_3 alkoxy, and -NR_{1-a}R_{1-b},

(iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, 10 -OH, -SH, -C≡N, -CF₃, C_1-C_3 alkoxy, and -NR_{1-a}R_{1-b},

(iv) -F, -Cl, -Br and -I,

(v) C_1-C_6 alkoxy optionally substituted with one, two, or three -F,

15 (vi) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,

(vii) -OH,

(viii) -C≡N,

(ix) $(CH_2)_{0-4}-C_3-C_7$ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, 20 -OH, -SH, -C≡N, -CF₃, C_1-C_3 alkoxy, and -NR_{1-a}R_{1-b},

(x) $(CH_2)_{0-4}-CO-(C_1-C_6 \text{ alkyl})$,

(xi) $(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,

(xii) $(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,

25 (xiii) $(CH_2)_{0-4}-SO_2-(C_1-C_6 \text{ alkyl})$,

(xiv) $(CH_2)_{0-4}-N(R_{N-2})-SO_2-$,

and

(xv) $(CH_2)_{0-4}-N(R_{N-2})-C(O)-$,

30 (8) $-(CH_2)_{0-4}-CO-(C_1-C_{12} \text{ alkyl})$,

(9) $-(CH_2)_{0-4}-CO-(C_2-C_{12} \text{ alkenyl})$,

(10) $-(CH_2)_{0-4}-CO-(C_2-C_{12} \text{ alkynyl})$,

(11) $-(CH_2)_{0-4}-CO-(CH_2)_{0-4} (C_3-C_7 \text{ cycloalkyl})$,

(12) $-(CH_2)_{0-4}-CO-R_{1-aryl}$,

(13) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl},$

(14) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle}$ wherein

$\text{R}_1\text{-heterocycle}$ at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl, dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl, dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the $\text{R}_1\text{-heterocycle}$ group is bonded by any atom of the parent $\text{R}_1\text{-heterocycle}$ group substituted by hydrogen such that the new bond to the $\text{R}_1\text{-heterocycle}$ group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(a) $\text{C}_1\text{-C}_6$ alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of $\text{C}_1\text{-C}_3$ alkyl, halogen, $-\text{OH}$, $-\text{SH}$, $-\text{NR}_{1-a}\text{R}_{1-b}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, and $\text{C}_1\text{-C}_3$ alkoxy,

(b) $\text{C}_2\text{-C}_6$ alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1\text{-C}_3$ alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$

(c) $\text{C}_2\text{-C}_6$ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1\text{-C}_3$ alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$

(d) halogen,

(e) $\text{C}_1\text{-C}_6$ alkoxy,

(f) $-\text{C}_1\text{-C}_6$ alkoxy optionally substituted with one, two, or three $-\text{F}$,

(g) $-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,(h) $-\text{OH}$,(i) $-\text{C}\equiv\text{N}$,(j) $(\text{CH}_2)_{0-4}-(\text{C}_3-\text{C}_7 \text{ cycloalkyl})$,

5 optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3 alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(k) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_4 \text{ alkyl})$,(l) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{1-a}\text{R}_{1-b}$,

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(m) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$,(n) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_6 \text{ alkyl})$, and(o) $=\text{O}$,(p) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{SO}_2-$ (q) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{C}(\text{O})-$

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(15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$ wherein

$\text{R}_{\text{N}-4}$ at each occurrence is independently selected from the group consisting of morpholinyl, thiomorpholinyl, pyrrolidinonyl, pyrrolyl, pyrazolyl, thienyl, pyridyl N-oxide, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl and pyrrolidinyl where each group is optionally substituted with 1, 2, 3, or 4 groups that are independently C_1-C_6 alkyl,

(16) $-(\text{CH}_2)_{0-4}-\text{CO}_2-\text{R}_{\text{N}-5}$ where

25

$\text{R}_{\text{N}-5}$ at each occurrence is independently selected from the group consisting of:

(a) C_1-C_6 alkyl,(b) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-aryl})$,(c) C_2-C_6 alkenyl,

30

(d) C_2-C_6 alkynyl,(e) C_3-C_7 cycloalkyl, and(f) $-(\text{CH}_2)_{0-4}-(\text{R}_1\text{-heteroaryl})$,(17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$ (18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl})$,

35

(19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,

- (20) - (CH₂)₀₋₄-SO₂-(C₃-C₇ cycloalkyl),
 (21) - (CH₂)₀₋₄-N(H or R_{N-5})-CO₂-R_{N-5},
 (22) - (CH₂)₀₋₄-N(H or R_{N-5})-CO-N(R_{N-5})₂,
 (23) - (CH₂)₀₋₄-N-CS-N(R_{N-5})₂,
 5 (24) - (CH₂)₀₋₄-N(-H or R_{N-5})-CO-R_{N-2},
 (25) - (CH₂)₀₋₄-NR_{N-2}R_{N-3},
 (26) - (CH₂)₀₋₄-R_{N-4},
 (27) - (CH₂)₀₋₄-O-CO-(C₁-C₆ alkyl),
 (28) - (CH₂)₀₋₄-O-P(O)-(OR₁₀₀)₂ where R₁₀₀ is
 10 independently H or C₁-C₄ alkyl,
 (29) - (CH₂)₀₋₄-O-CO-N(R_{N-5})₂,
 (30) - (CH₂)₀₋₄-O-CS-N(R_{N-5})₂,
 (31) - (CH₂)₀₋₄-O-(R_{N-5}),
 (32) - (CH₂)₀₋₄-O-(R_{N-5})-COOH,
 15 (33) - (CH₂)₀₋₄-S-(R_{N-5}),
 (34) - (CH₂)₀₋₄-O-(C₁-C₆ alkyl) wherein the
 alkyl group is optionally substituted with one, two, three,
 four, or five substituents independently selected from the group
 consisting of F, Cl, Br, and I,
 20 (35) - (CH₂)₀₋₄-(C₃-C₈ cycloalkyl),
 (36) C₂-C₆ alkenyl optionally substituted
 with C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, or
 -NR_{1-a}R_{1-b},
 (37) C₂-C₆ alkynyl optionally substituted
 25 with C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃
 alkoxy, or -NR_{1-a}R_{1-b}, and
 (38) - (CH₂)₀₋₄-N(-H or R_{N-5})-SO₂-R_{N-2};
 (IV) - (CR_{C-x}R_{C-y})₀₋₄-R_{C-heteroaryl} wherein R_{C-heteroaryl} at each
 occurrence is independently selected from the group consisting
 30 of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl,
 indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl,
 quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl,
 pyrazolyl, oxazolyl, thiazolyl, indolizinyl, indazolyl,
 benzoisothiazolyl, benzimidazolyl, benzofuranyl, furanyl,
 35 thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl,

tetrazolyl, oxazolopyridinyl, isothiazolyl, naphthyridinyl,
 cinnolinyl, carbazolyl, beta-carbolinyl, isochromanlyl, chromanlyl,
 tetrahydroisoquinolinyl, isoindolinyl,
 isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl,
 5 isobenzothienyl, benzoxazolyl, pyridopyridinyl,
 benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl,
 benzodioxolyl, triazinyl, henoxazinyl, phenothiazinyl,
 pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
 dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
 10 dihydrobenzisothiazinyl, benzopyranlyl, benzothiopyranlyl,
 coumarinyl, isocoumarinyl, chromonyl, chromanonyl,
 tetrahydroquinolinyl, dihydroquinolinyl,
 dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
 dihydroisocoumarinyl, isoindolinonyl,
 15 benzodioxanlyl, benzoxazolinonyl, imidazopyrazolyl,
 quinazolinonyl, pyrazopyridyl, benzooxadiazolyl,
 dihydropyrimidinonyl, dihydrobenzofuranonyl,

where the R_C-heteroaryl group is bonded by any atom of the
 parent R_C-heteroaryl group substituted by hydrogen such that the new
 20 bond to the R_C-heteroaryl group replaces the hydrogen atom and its
 bond, where heteroaryl is optionally substituted 1, 2, 3, or 4
 groups that are independently:

(1) C₁-C₆ alkyl, optionally substituted with 1, 2, or
 3 groups independently selected from the group consisting of C₁-
 25 C₃ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy,
 and -NR_{1-a}R_{1-b},

(2) -OH,

(3) -NO₂,

(4) -F, -Cl, -Br, -I,

30 (5) -CO-OH,

(6) -C≡N,

(V) C₂-C₁₀ alkenyl optionally substituted with one, two
 or three substituents independently selected from the group

consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_6 alkoxy, -O-phenyl, and -NR_{1-a}R_{1-b},

(VI) C_2 - C_{10} alkynyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_6 alkoxy, -O-phenyl, and -NR_{1-a}R_{1-b},

(VII) -(C_1 - C_6 alkyl)-O-(C_1 - C_6 alkyl)-OH,

(VIII) -CH₂-NH-CH₂-CH(-O-CH₂-CH₃)₂,

(IX) -(CH₂)₀₋₆-C(=NR_{1-a})(NR_{1-a}R_{1-b});

10 where R_N is

(I) R_{N-1}-X_N- where X_N is -CO-, and where R_{N-1} is selected from the group consisting of:

(A) phenyl, which is optionally substituted with one, two or three of the following substituents which can be the same or different and are:

(1) C_1 - C_6 alkyl, optionally substituted with one, two or three substituents selected from the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},

20 wherein R_{1-a} and R_{1-b} at each occurrence are independently H or C_1 - C_6 alkyl,

(2) -OH,

(3) -NO₂,

(4) -F, -Cl, -Br, -I,

25 (5) -CO₂H,

(6) -C \equiv N,

(7) -(CH₂)₀₋₄-CO-NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are the same or different and are selected from the group consisting of:

(a) -H,

30 (b) C_1 - C_8 alkyl optionally substituted with one substituent selected from the group consisting of:

(i) -OH,

(ii) -NH₂,

(iii) phenyl,

(c) $-C_1-C_8$ alkyl optionally substituted with 1, 2, or 3 groups that are independently $-F$, $-Cl$, $-Br$, or $-I$,

(d) $-C_3-C_8$ cycloalkyl,

(e) $-(C_1-C_2 \text{ alkyl})-(C_3-C_8 \text{ cycloalkyl})$,

5 (f) $-(C_1-C_6 \text{ alkyl})-O-(C_1-C_3 \text{ alkyl})$,

(g) $-C_2-C_6$ alkenyl,

(h) $-C_2-C_6$ alkynyl,

(i) $-C_1-C_6$ alkyl chain with one double bond and one triple bond,

10 (j) $-R_{1-aryl}$, wherein R_{1-aryl} at each occurrence is independently phenyl, naphthyl, indanyl, indenyl, dihydronaphthyl, or tetralinyl each of which is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(i) C_1-C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

(ii) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

20 (iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of $-F$, $-Cl$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(iv) $-F$, Cl , $-Br$ and $-I$,

(v) $-C_1-C_6$ alkoxy optionally substituted with 1, 2, or 3 $-F$,

(vi) $-NR_{N-2}R_{N-3}$,

30 (vii) $-OH$,

(viii) $-C\equiv N$,

(ix) C_3-C_7 cycloalkyl, optionally substituted with 1, 2, or 3 groups that are selected from the

group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(x) -CO-(C₁-C₄ alkyl),

(xi) -SO₂-NR_{1-a}R_{1-b},

5 (xii) -CO-NR_{1-a}R_{1-b}, or

(xiii) -SO₂-(C₁-C₄ alkyl),

(k) -R₁-heteroaryl, wherein R₁-heteroaryl at each occurrence is independently selected from the group consisting of pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, 10 indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl, quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indoliziny, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, 15 triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl, isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, 20 pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl, purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl, pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl, dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl, dihydrobenzisothiazinyl, benzopyranyl, benzothiopyranyl, 25 coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-oxide, tetrahydroquinolinyl, dihydroquinolinyl, dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl, dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl, benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide, 30 pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide, indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide, quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide, thiazolyl N-oxide, indoliziny, indazolyl N-oxide, 35 benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-

oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide, and benzothiopyranyl S,S-dioxide,

where the R_1 -heteroaryl group is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(i) C_1-C_6 alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of C_1-C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

(ii) C_2-C_6 alkenyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, or $-NR_{1-a}R_{1-b}$,

(iii) C_2-C_6 alkynyl optionally substituted with 1, 2, or 3 groups that are independently -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, or $-NR_{1-a}R_{1-b}$,

(iv) -F, -Cl, -Br and -I,

(v) $-C_1-C_6$ alkoxy optionally substituted with one, two, or three -F,

(vi) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,

(vii) -OH,

(viii) $-C\equiv N$,

(ix) $(CH_2)_{0-4}-C_3-C_7$ cycloalkyl, optionally substituted with 1, 2, or 3 groups that are independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$,

(x) $(CH_2)_{0-4}-CO-(C_1-C_6 \text{ alkyl})$,

(xi) $(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,

(xii) $(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,

(xiii) $(CH_2)_{0-4}-SO_2-(C_1-C_6 \text{ alkyl})$,

(xiv) $(CH_2)_{0-4}-N(R_{N-2})-SO_2-$, and

(xv) $(CH_2)_{0-4}-N(R_{N-2})-C(O)-$,

(l) $-R_1$ -heterocycle, wherein

R_1 -heterocycle at each occurrence is independently selected from the group consisting of morpholinyl,

thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl, dihydropyrazinyl, dihydropyridinyl, dihydropyrimidinyl, dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, and homothiomorpholinyl S-oxide,

where the R_1 -heterocycle group is bonded by any atom of the parent R_1 -heterocycle group substituted by hydrogen such that the new bond to the R_1 -heterocycle group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with 1, 2, 3, or 4 groups that are independently:

(a) C_1-C_6 alkyl optionally substituted with one, two or three substituents independently selected from the group consisting of C_1-C_3 alkyl, halogen, -OH, -SH, $-NR_{1-a}R_{1-b}$, $-C\equiv N$, $-CF_3$, and C_1-C_3 alkoxy,

(b) C_2-C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(c) C_2-C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents independently selected from the group consisting of -F, -Cl, -OH, -SH, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, and $-NR_{1-a}R_{1-b}$

(d) halogen,

(e) C_1-C_6 alkoxy,

(f) $-C_1-C_6$ alkoxy optionally substituted with one, two, or three -F,

(g) $-NR_{N-2}R_{N-3}$,

(h) -OH,

(i) $-C\equiv N$,

(j) $(\text{CH}_2)_{0-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$, optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

- 5 (k) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_4 \text{ alkyl})$,
(l) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{1-a}\text{R}_{1-b}$,
(m) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$,
(n) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_6 \text{ alkyl})$, and
(o) =O,
10 (p) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{SO}_2-$
(q) $-(\text{CH}_2)_{0-4}-\text{N}(\text{R}_{\text{N}-2})-\text{C}(\text{O})-$

- (8) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
(9) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkenyl})$,
(10) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_2-\text{C}_{12} \text{ alkynyl})$,
15 (11) $-(\text{CH}_2)_{0-4}-\text{CO}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
(12) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-aryl}$,
(13) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heteroaryl}$,
(14) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_1\text{-heterocycle}$,
(15) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{R}_{\text{N}-4}$ wherein R_{N-4} is selected from
20 the group consisting of phenyl, morpholinyl, thiomorpholinyl, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl, thienyl, pyrazolyl, pyridyl N-oxide, oxazolyl, thiazolyl, imidazolyl, and pyrrolidinyl where each group is
25 optionally substituted with one, two, three, or four groups that are independently C₁-C₆ alkyl,

(16) $-(\text{CH}_2)_{0-4}-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$ where R_{N-5} is selected from the group consisting of:

- (a) C₁-C₆ alkyl,
30 (b) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-aryl})$,
(c) C₂-C₆ alkenyl,
(d) C₂-C₆ alkynyl,
(e) $-(\text{CH}_2)_{0-2}-\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,
(f) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heteroaryl})$, and
35 (g) $-(\text{CH}_2)_{0-2}-(\text{R}_1\text{-heterocycle})$,

- (17) $-(\text{CH}_2)_{0-4}-\text{SO}_2-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
 (18) $-(\text{CH}_2)_{0-4}-\text{SO}-(\text{C}_1-\text{C}_8 \text{ alkyl})$,
 (19) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_1-\text{C}_{12} \text{ alkyl})$,
 (20) $-(\text{CH}_2)_{0-4}-\text{SO}_2-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,
 5 (21) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{O}-\text{R}_{\text{N}-5}$,
 (22) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (23) $-(\text{CH}_2)_{0-4}-\text{N}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (24) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{CO}-\text{R}_{\text{N}-2}$,
 (25) $-(\text{CH}_2)_{0-4}-\text{NR}_{\text{N}-2}\text{R}_{\text{N}-3}$,
 10 (26) $-(\text{CH}_2)_{0-4}-\text{R}_{\text{N}-4}$,
 (27) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,
 (28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$ wherein
 R_{100} at each occurrence is independently -H
 or $\text{C}_1-\text{C}_4 \text{ alkyl}$,
 15 (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
 (32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
 (33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
 20 (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl optionally}$
 substituted with one, two, three, four, or five of -F),
 (35) $\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,
 (36) $\text{C}_2-\text{C}_6 \text{ alkenyl optionally substituted with } \text{C}_1-$
 $\text{C}_3 \text{ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C}\equiv\text{N, -CF}_3, \text{C}_1-\text{C}_3 \text{ alkoxy,}$
 25 or $-\text{NR}_{1-a}\text{R}_{1-b}$,
 (37) $\text{C}_2-\text{C}_6 \text{ alkynyl optionally substituted with } \text{C}_1-$
 $\text{C}_3 \text{ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C}\equiv\text{N, -CF}_3, \text{C}_1-\text{C}_3 \text{ alkoxy,}$
 or $-\text{NR}_{1-a}\text{R}_{1-b}$,
 30 (38) $-(\text{CH}_2)_{0-4}-\text{N}(\text{H or } \text{R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$,
 (39) $-(\text{CH}_2)_{1-4}-(\text{C}_3-\text{C}_8 \text{ cycloalkyl})$,

(B) $-\text{R}_{\text{N-heteroaryl}}$ where $\text{R}_{\text{N-heteroaryl}}$ is selected from the group consisting of pyridinyl, indolyl, indolinyl, isoindolyl, imidazolyl, isoxazolyl, oxazolyl, thiazolyl, indolizinyll and isochromanyl,

where the $R_{N\text{-heteroaryl}}$ group is bonded by any atom of the parent $R_{N\text{-heteroaryl}}$ group substituted by hydrogen such that the new bond to the $R_{N\text{-heteroaryl}}$ group replaces the hydrogen atom and its bond, where heteroaryl is optionally substituted with one, two, three, or four of:

(1) $C_1\text{-}C_6$ alkyl, optionally substituted with one, two or three substituents selected from the group consisting of $C_1\text{-}C_3$ alkyl, $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, $C_1\text{-}C_3$ alkoxy, and $-NR_{1-a}R_{1-b}$,

- (2) $-OH$,
- (3) $-NO_2$,
- (4) $-F$, $-Cl$, $-Br$, $-I$,
- (5) $-CO_2H$,
- (6) $-C\equiv N$,
- (7) $-(CH_2)_{0-4}-CO-NR_{N-2}R_{N-3}$,
- (8) $-(CH_2)_{0-4}-CO-(C_1\text{-}C_{12} \text{ alkyl})$,
- (9) $-(CH_2)_{0-4}-CO-(C_2\text{-}C_{12} \text{ alkenyl})$,
- (10) $-(CH_2)_{0-4}-CO-(C_2\text{-}C_{12} \text{ alkynyl})$,
- (11) $-(CH_2)_{0-4}-CO-(C_3\text{-}C_8 \text{ cycloalkyl})$,
- (12) $-(CH_2)_{0-4}-CO-R_{1\text{-aryl}}$,
- (13) $-(CH_2)_{0-4}-CO-R_{1\text{-heteroaryl}}$,
- (14) $-(CH_2)_{0-4}-CO-R_{1\text{-heterocycle}}$,
- (15) $-(CH_2)_{0-4}-CO-R_{N-4}$,
- (16) $-(CH_2)_{0-4}-CO-O-R_{N-5}$,
- (17) $-(CH_2)_{0-4}-SO_2-NR_{N-2}R_{N-3}$,
- (18) $-(CH_2)_{0-4}-SO-(C_1\text{-}C_8 \text{ alkyl})$,
- (19) $-(CH_2)_{0-4}-SO_2-(C_1\text{-}C_{12} \text{ alkyl})$,
- (20) $-(CH_2)_{0-4}-SO_2-(C_3\text{-}C_8 \text{ cycloalkyl})$,
- (21) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO-O-R_{N-5}$,
- (22) $-(CH_2)_{0-4}-N(H \text{ or } R_{N-5})-CO-N(R_{N-5})_2$,
- (23) $-(CH_2)_{0-4}-N-CS-N(R_{N-5})_2$,
- (24) $-(CH_2)_{0-4}-N(-H \text{ or } R_{N-5})-CO-R_{N-2}$,
- (25) $-(CH_2)_{0-4}-NR_{N-2}R_{N-3}$,
- (26) $-(CH_2)_{0-4}-R_{N-4}$,
- (27) $-(CH_2)_{0-4}-O-CO-(C_1\text{-}C_6 \text{ alkyl})$,

- (28) $-(\text{CH}_2)_{0-4}-\text{O}-\text{P}(\text{O})-(\text{OR}_{100})_2$,
 (29) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CO}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (30) $-(\text{CH}_2)_{0-4}-\text{O}-\text{CS}-\text{N}(\text{R}_{\text{N}-5})_2$,
 (31) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})$,
 5 (32) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{R}_{\text{N}-5})-\text{COOH}$,
 (33) $-(\text{CH}_2)_{0-4}-\text{S}-(\text{R}_{\text{N}-5})$,
 (34) $-(\text{CH}_2)_{0-4}-\text{O}-(\text{C}_1-\text{C}_6 \text{ alkyl optionally substituted with one, two, three, four, or five of -F})$,
 (35) $\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,
 10 (36) $\text{C}_2-\text{C}_6 \text{ alkenyl optionally substituted with } \text{C}_1-\text{C}_3 \text{ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C}\equiv\text{N, -CF}_3, \text{C}_1-\text{C}_3 \text{ alkoxy, or -NR}_{1-\text{a}}\text{R}_{1-\text{b}}$,
 (37) $\text{C}_2-\text{C}_6 \text{ alkynyl optionally substituted with } \text{C}_1-\text{C}_3 \text{ alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C}\equiv\text{N, -CF}_3, \text{C}_1-\text{C}_3 \text{ alkoxy, or -NR}_{1-\text{a}}\text{R}_{1-\text{b}}$,
 15 (38) $-(\text{CH}_2)_{0-4}-\text{N}(-\text{H or R}_{\text{N}-5})-\text{SO}_2-\text{R}_{\text{N}-2}$,
 (39) $-(\text{CH}_2)_{1-4}-\text{C}_3-\text{C}_8 \text{ cycloalkyl}$,
 (C) $\text{R}_{\text{N-aryl}}-\text{W}-\text{R}_{\text{N-aryl}}$,
 (D) $\text{R}_{\text{N-aryl}}-\text{W}-\text{R}_{\text{N-heteroaryl}}$,
 20 (E) $\text{R}_{\text{N-aryl}}-\text{W}-\text{R}_{1-\text{heterocycle}}$,
 (F) $\text{R}_{\text{N-heteroaryl}}-\text{W}-\text{R}_{\text{N-aryl}}$,
 (G) $\text{R}_{\text{N-heteroaryl}}-\text{W}-\text{R}_{\text{N-heteroaryl}}$,
 (H) $\text{R}_{\text{N-heteroaryl}}-\text{W}-\text{R}_{\text{N-1-heterocycle}}$,
 (I) $\text{R}_{\text{N-heterocycle}}-\text{W}-\text{R}_{\text{N-aryl}}$,
 25 (J) $\text{R}_{\text{N-heterocycle}}-\text{W}-\text{R}_{\text{N-heteroaryl}}$,
 (K) $\text{R}_{\text{N-heterocycle}}-\text{W}-\text{R}_{\text{N-1-heterocycle}}$,

where W is

- (25) $-(\text{CH}_2)_{1-4}-$,
 (26) $-\text{O}-$,
 30 (27) $-\text{S}(\text{O})_{0-2}-$,
 (28) $-\text{N}(\text{R}_{\text{N}-5})-$,
 (29) $-\text{CO}-$; or
 (30) a bond;

(II) $-\text{CO}-(\text{C}_1-\text{C}_6 \text{ alkyl})-\text{M}-(\text{C}_1-\text{C}_6 \text{ alkyl})$, where M is S, SO or
 35 SO_2 , and wherein each alkyl is unsubstituted or substituted with

one, two, or three of substituents independently selected from the group consisting of:

(A) $\text{-NH-CO-(C}_1\text{-C}_6\text{ alkyl)}$,

(B) -NH-CO-O-R_{N-8} ,

5 (C) $\text{-NR}_{N-2}\text{R}_{N-3}$;

where R_1 is

$\text{-(CH}_2\text{)}_{n_1}\text{-phenyl}$, where n_1 is zero or one, and which is optionally substituted with one, two, three or four of the following substituents on the phenyl ring:

10 (A) $\text{C}_1\text{-C}_6$ alkyl optionally substituted with one, two or three substituents selected from the group consisting of $\text{C}_1\text{-C}_3$ alkyl, -F , -Cl , -Br , -I , -OH , -SH , $\text{-C}\equiv\text{N}$, -CF_3 , $\text{C}_1\text{-C}_3$ alkoxy, and $\text{-NR}_{1-a}\text{R}_{1-b}$,

(B) $\text{C}_2\text{-C}_6$ alkenyl with one or two double bonds, 15 optionally substituted with one, two or three substituents selected from the group consisting of -F , -Cl , -OH , -SH , $\text{-C}\equiv\text{N}$, -CF_3 , $\text{C}_1\text{-C}_3$ alkoxy, and $\text{-NR}_{1-a}\text{R}_{1-b}$,

(C) $\text{C}_2\text{-C}_6$ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents 20 selected from the group consisting of -F , -Cl , -OH , -SH , $\text{-C}\equiv\text{N}$, -CF_3 , $\text{C}_1\text{-C}_3$ alkoxy, and $\text{-NR}_{1-a}\text{R}_{1-b}$,

(D) -F , Cl , -Br or -I ,

(F) $\text{-C}_1\text{-C}_6$ alkoxy optionally substituted with one, two or three of -F ,

25 (G) $\text{-NR}_{N-2}\text{R}_{N-3}$ where R_{N-2} and R_{N-3} are as defined below,

(H) -OH ,

(I) $\text{-C}\equiv\text{N}$,

(J) $\text{C}_3\text{-C}_7$ cycloalkyl, optionally substituted with 30 one, two or three substituents selected from the group consisting of -F , -Cl , -OH , -SH , $\text{-C}\equiv\text{N}$, -CF_3 , $\text{C}_1\text{-C}_3$ alkoxy, and $\text{-NR}_{1-a}\text{R}_{1-b}$,

(K) $\text{-CO-(C}_1\text{-C}_4\text{ alkyl)}$,

(L) $\text{-SO}_2\text{-NR}_{1-a}\text{R}_{1-b}$,

(M) $-\text{CO}-\text{NR}_{1-a}\text{R}_{1-b}$,

(N) $-\text{SO}_2-(\text{C}_1-\text{C}_4 \text{ alkyl})$; and

where R_2 is

(I) $-(\text{Z})-\text{C}_1-\text{C}_6 \text{ alkyl}$, where Z is a bond, $-\text{C}(\text{O})-$, $-\text{CO}_2-$
5 or $-\text{SO}_2-$, wherein the alkyl group is optionally substituted with
one, two or three substituents selected from the group
consisting of $\text{C}_1-\text{C}_3 \text{ alkyl}$, $\text{C}_1-\text{C}_7 \text{ alkyl}$ (optionally substituted
with $\text{C}_1-\text{C}_3 \text{ alkyl}$ and $\text{C}_1-\text{C}_3 \text{ alkoxy}$), $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$,
 $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, $-\text{NR}_{1-a}\text{R}_{1-b}$ where R_{1-a} and R_{1-b} are
10 independently $-\text{H}$ or $\text{C}_1-\text{C}_6 \text{ alkyl}$, and $-\text{OC}=\text{O NR}_{1-a}\text{R}_{1-b}$,

(II) $-(\text{Z})-\text{CH}_2-\text{S}(\text{O})_{0-2}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,

(III) $-(\text{Z})-\text{CH}_2-\text{CH}_2-\text{S}(\text{O})_{0-2}-(\text{C}_1-\text{C}_6 \text{ alkyl})$,

(IV) $-(\text{Z})-\text{C}_2-\text{C}_6 \text{ alkenyl}$ with one or two double bonds,
optionally substituted with one, two or three substituents
15 selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(V) $-(\text{Z})-\text{C}_2-\text{C}_6 \text{ alkynyl}$ with one or two triple bonds,
optionally substituted with one, two or three substituents
selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, and $-\text{NR}_{1-a}\text{R}_{1-b}$,
20

(VI) $-(\text{Z})-(\text{CH}_2)_{n_1}-(\text{R}_{1-\text{aryl}})$, where Z is a bond, CO, CO_2
or SO_2 , where n_1 is zero or one and where $\text{R}_{1-\text{aryl}}$ is phenyl, 1-
naphthyl, 2-naphthyl and indanyl, indenyl, dihydronaphthalenyl, or
tetralinyl optionally substituted with one, two, three or four
25 of the following substituents on the aryl ring:

(A) $\text{C}_1-\text{C}_6 \text{ alkyl}$ optionally substituted with one,
two or three substituents selected from the group consisting of
 $\text{C}_1-\text{C}_3 \text{ alkyl}$, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, C_1-C_3
alkoxy, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

30 (B) $\text{C}_2-\text{C}_6 \text{ alkenyl}$ with one or two double bonds,
optionally substituted with one, two or three substituents
selected from the group consisting of $-\text{F}$, $-\text{Cl}$, $-\text{OH}$, $-\text{SH}$, $-\text{C}\equiv\text{N}$, $-\text{CF}_3$, $\text{C}_1-\text{C}_3 \text{ alkoxy}$, and $-\text{NR}_{1-a}\text{R}_{1-b}$,

(C) C₂-C₆ alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

5 (D) -F, Cl, -Br or -I,

(F) -C₁-C₆ alkoxy optionally substituted with one, two or three of - F,

(G) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

10 (H) -OH,

(I) -C≡N,

(J) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -
15 NR_{1-a}R_{1-b},

(K) -CO-(C₁-C₄ alkyl),

(L) -SO₂-NR_{1-a}R_{1-b},

(M) -CO-NR_{1-a}R_{1-b},

(N) -SO₂-(C₁-C₄ alkyl),

20 (VII) -(Z)-(CH₂)_{n1}-(R_{1-heteroaryl}) where n₁ is as defined above and where R_{1-heteroaryl} is selected from the group consisting of:

pyridinyl, pyrimidinyl, quinolinyl, benzothienyl, indolyl, indolinyl, pyridazinyl, pyrazinyl, isoindolyl, isoquinolyl,
25 quinazolinyl, quinoxalinyl, phthalazinyl, imidazolyl, isoxazolyl, pyrazolyl, oxazolyl, thiazolyl, indolizinyl, indazolyl, benzothiazolyl, benzimidazolyl, benzofuranyl, furanyl, thienyl, pyrrolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, oxazolopyridinyl, imidazopyridinyl,
30 isothiazolyl, naphthyridinyl, cinnolinyl, carbazolyl, beta-carbolinyl, isochromanyl, chromanyl, tetrahydroisoquinolinyl, isoindolinyl, isobenzotetrahydrofuranlyl, isobenzotetrahydrothienyl, isobenzothienyl, benzoxazolyl, pyridopyridinyl, benzotetrahydrofuranlyl, benzotetrahydrothienyl,

purinyl, benzodioxolyl, triazinyl, phenoxazinyl, phenothiazinyl,
 pteridinyl, benzothiazolyl, imidazopyridinyl, imidazothiazolyl,
 dihydrobenzisoxazinyl, benzisoxazinyl, benzoxazinyl,
 dihydrobenzisoctiazinyl, benzopyranyl, benzothiopyranyl,
 5 coumarinyl, isocoumarinyl, chromonyl, chromanonyl, pyridinyl-N-
 oxide, tetrahydroquinolinyl, dihydroquinolinyl,
 dihydroquinolinonyl, dihydroisoquinolinonyl, dihydrocoumarinyl,
 dihydroisocoumarinyl, isoindolinonyl, benzodioxanyl,
 benzoxazolinonyl, pyrrolyl N-oxide, pyrimidinyl N-oxide,
 10 pyridazinyl N-oxide, pyrazinyl N-oxide, quinolinyl N-oxide,
 indolyl N-oxide, indolinyl N-oxide, isoquinolyl N-oxide,
 quinazolinyl N-oxide, quinoxalinyl N-oxide, phthalazinyl N-
 oxide, imidazolyl N-oxide, isoxazolyl N-oxide, oxazolyl N-oxide,
 thiazolyl N-oxide, indolizinyll N-oxide, indazolyl N-oxide,
 15 benzothiazolyl N-oxide, benzimidazolyl N-oxide, pyrrolyl N-
 oxide, oxadiazolyl N-oxide, thiadiazolyl N-oxide, triazolyl N-
 oxide, tetrazolyl N-oxide, benzothiopyranyl S-oxide,
 benzothiopyranyl S,S-dioxide,

where the R₁-heteroaryl group is bonded to -(CH₂)_{n1}- by any ring atom
 20 of the parent R_N-heteroaryl group substituted by hydrogen such that
 the new bond to the R₁-heteroaryl group replaces the hydrogen atom
 and its bond, where heteroaryl is optionally substituted with
 one, two, three or four of:

(1) C₁-C₆ alkyl optionally substituted with one,
 25 two or three substituents selected from the group consisting of
 C₁-C₃ alkyl, -F, -Cl, -Br, -I, -OH,
 -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(2) C₂-C₆ alkenyl with one or two double bonds,
 optionally substituted with one, two or three substituents
 30 selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -
 CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(3) C₂-C₆ alkynyl with one or two triple bonds,
 optionally substituted with one, two or three substituents

selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(4) -F, Cl, -Br or -I,

(6) -C₁-C₆ alkoxy optionally substituted with one, two, or three of -F,

(7) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,

(8) -OH,

(9) -C≡N,

(10) C₃-C₇ cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b},

(11) -CO-(C₁-C₄ alkyl),

(12) -SO₂-NR_{1-a}R_{1-b},

(13) -CO-NR_{1-a}R_{1-b}, or

(14) -SO₂-(C₁-C₄ alkyl), with the proviso that when n₁ is zero R_{1-heteroaryl} is not bonded to the carbon chain by nitrogen, or

(VIII) -(Z)-(CH₂)_{n1}-(R_{1-heterocycle}) where n₁ is as defined above and R_{1-heterocycle} is selected from the group consisting of:

morpholinyl, thiomorpholinyl, thiomorpholinyl S-oxide, thiomorpholinyl S,S-dioxide, piperazinyl, homopiperazinyl, pyrrolidinyl, pyrrolinyl, tetrahydropyranyl, piperidinyl, tetrahydrofuranyl, tetrahydrothienyl, homopiperidinyl, homomorpholinyl, homomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, oxazolidinonyl, dihydropyrazolyl, dihydropyrrolyl dihydropyrazinyl dihydropyridinyl dihydropyrimidinyl, dihydrofuryl, dihydropyranyl, tetrahydrothienyl S-oxide, tetrahydrothienyl S,S-dioxide, homothiomorpholinyl S-oxide,

where the R_{1-heterocycle} group is bonded by any atom of the parent R_{1-heterocycle} group substituted by hydrogen such that the new bond

to the R_1 -heterocycle group replaces the hydrogen atom and its bond, where heterocycle is optionally substituted with one, two, three or four:

- (1) C_1 - C_6 alkyl optionally substituted with one, two or three substituents selected from the group consisting of C_1 - C_3 alkyl, -F, -Cl, -Br, -I, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},
- (2) C_2 - C_6 alkenyl with one or two double bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},
- (3) C_2 - C_6 alkynyl with one or two triple bonds, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},
- (4) -F, Cl, -Br, or -I,
- (5) C_1 - C_6 alkoxy,
- (6) - C_1 - C_6 alkoxy optionally substituted with one, two, or three -F,
- (7) -NR_{N-2}R_{N-3} where R_{N-2} and R_{N-3} are as defined below,
- (8) -OH,
- (9) -C \equiv N,
- (10) C_3 - C_7 cycloalkyl, optionally substituted with one, two or three substituents selected from the group consisting of -F, -Cl, -OH, -SH, -C \equiv N, -CF₃, C_1 - C_3 alkoxy, and -NR_{1-a}R_{1-b},
- (11) -CO-(C_1 - C_4 alkyl),
- (12) -SO₂-NR_{1-a}R_{1-b},
- (13) -CO-NR_{1-a}R_{1-b},
- (14) -SO₂-(C_1 - C_4 alkyl),
- (15) =O, with the proviso that when n_1 is zero R_1 -heterocycle is not bonded to the carbon chain by nitrogen; and

where R_{20} is H or C_{1-6} alkyl or alkenyl.

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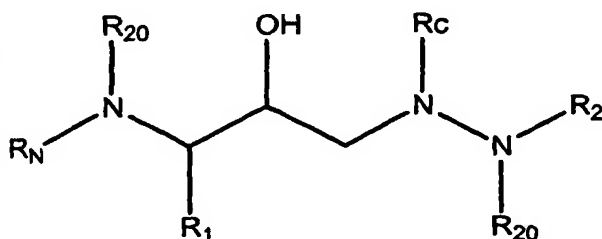
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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: AZA HYDROXYLATED ETHYL AMINE COMPOUNDS



(I)

(57) Abstract: Disclosed are compounds of formula (I) and pharmaceutically acceptable salts and esters thereof, useful in treating and/or preventing Alzheimer's disease and other similar diseases, wherein R_N , R_C , R_1 , R_2 and R_{20} are defined herein. These compounds include inhibitors of the beta-secretase enzyme that are useful in the treatment of Alzheimer's disease and other diseases characterized by deposition of A beta peptide in a mammal. The compounds of the invention are useful in pharmaceutical compositions and methods of treatment to reduce A beta peptide formation.

WO 02/094768 A3

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 02/16199

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07C243/28 A61K31/16 A61P25/28 C07D277/56

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, BEILSTEIN Data, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 02 02506 A (ELAN PHARM INC) 10 January 2002 (2002-01-10) page 1, line 14 - line 29 page 106 -page 107; examples 1-3 chart A, B and C'	1-36
A	WO 99 36404 A (SQUIBB BRISTOL MYERS CO) 22 July 1999 (1999-07-22) page 1, line 1 - line 20	1-36
A	US 6 225 345 B1 (BOLD GUIDO ET AL) 1 May 2001 (2001-05-01) column 2, line 6 - line 64	1-36

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the International filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the International filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 02/16199

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claim 36 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound.
2. ☒ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

see FURTHER INFORMATION sheet PCT/ISA/210
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Present claims 1-3, 36 relate to an extremely large number of possible compounds and their uses. In fact, the claims contain so many options, variables, possible permutations and run to such unnecessary length (23 pages for claim 1) that a lack of conciseness within the meaning of Article 6 PCT arises to such an extent as to render a meaningful search of the claims impossible. Consequently, the search has been carried out for those parts of the application which do appear to be concise namely the subject matter of claims 4-35. Claim 36 has also been searched with regard to the alleged effects of the compounds of claims 4-35.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

Information on patent family members

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